

GenCore version 5.1.6
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OK protein - protein search, using sw model

Run on: October 7, 2003, 18:46:24 ; Search time 19 seconds

(without alignments)
597.258 Million cell updates/sec

Title: US-09-898-860-2

Perfect score: 649

Sequence: 1 MPREDAHRTGYPKKGHSHS.....NAPPAYEKLSEOSPPPYSP 118

Scoring table:

BLOSUM62

Gap 10.0 , Gapext 0.5

Searched: 283308 seqs, 96168682 residues

Total number of hits satisfying chosen parameters: 283308

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

1: p1r1:*
2: p1r2:*
3: p1r3:*
4: p1r4:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	649	100.0	118	2 A55253	melanoma antigen M
2	87	13.4	4957	2 T03455	ALR protein - huma
3	87	13.4	5262	2 T03454	ALR protein - huma
4	80.5	12.4	344	2 B28967	T-cell surface gly
5	80.5	12.4	344	2 T49585	CD2 antigen protei
6	79	12.2	972	2 T49773	related to actin-I
7	77	11.9	710	2 E69665	nitrate reductase
8	73	11.2	1069	2 T00043	BH-protocadherin-a
9	72.5	11.2	1429	2 T13720	gene expanded prot
10	71.5	11.0	1265	2 T51314	probable CO-induce
11	71	10.9	1072	2 T00041	BH-protocadherin P
12	71	10.9	1200	2 T00042	BH-protocadherin P
13	70.5	10.9	354	2 H83334	probable transcript
14	70	10.8	335	2 G88640	protein F52C12.3 f
15	70	10.8	346	2 H84512	hypothetical prote
16	70	10.8	2100	2 T03323	probable polypeptid
17	69.5	10.7	341	2 T22533	hypothetical prote
18	69.5	10.7	404	2 H64175	hypothetical prote
19	69	10.6	940	2 S19702	fibronectin-blindi
20	68.5	10.6	384	2 E72100	hypothetical prote
21	68.5	10.6	476	2 T35769	probable transmem
22	68	10.5	316	2 C82085	conserved hypotet
23	68	10.5	377	2 C91270	beta-lactamase [im
24	68	10.5	377	2 C86111	beta-lactamase, pe
25	67.5	10.4	188	2 A64332	probable phosphono
26	67.5	10.4	315	2 T47971	seven in absentia-
27	67.5	10.4	476	2 T43464	hypothetical prote
28	67.5	10.4	670	2 T13739	probable hormone r
29	67.5	10.4	753	2 J00532	OP protein - Kenne

30	67	10.3	485	2 S32171	hydroxyneurosporen
31	67	10.3	2723	2 T03321	probable polyketid
32	66.5	10.2	215	2 F71923	hypothetical prote
33	66	10.2	140	2 S67666	probable membrane
34	66	10.2	751	2 T31515	hypothetical prote
35	65.5	10.1	258	2 A48820	homeobox protein (
36	65.5	10.1	384	2 A86521	conserved hypotet
37	65.5	10.1	518	2 D69539	antigen WCl.1 prec
38	65.5	10.1	1023	2 T48997	hypothetical prote
39	65.5	10.1	1436	2 A46496	epsin-like protein
40	65	10.0	309	1 E65112	hypothetical 34.6
41	65	10.0	309	2 B91140	hypothetical prote
42	65	10.0	309	2 E85985	conserved hypotet
43	65	10.0	309	2 AH0906	testicular protein
44	64.5	9.9	243	2 JEO204	fixz protein - Rhl
45	64.5	9.9	359	2 A22891	

ALIGNMENTS

RESULT 1
A55253
melanoma antigen MART-1 - human
N:Alternate names: melan-A protein
C:Species: Homo sapiens (man)
C>Date: 06-Feb-1995 #sequence_revision 06-Feb-1995 #text_change 04-Mar-2000
R:Kawakami, Y.; Elyahu, S.; Delgado, C.H.; Robbins, P.F.; Rivoltini, L.; Topalian
Proc. Natl. Acad. Sci. U.S.A. 91, 3515-3519, 1994
A>Title: Cloning of the gene coding for a shared human melanoma antigen recognized
A:Reference number: A55253; MUID:94224770; PMID:8170938
A:Accession: A55253
A>Status: preliminary
A:Molecule type: mRNA
A:Residues: 1-118 <KAM>
A:Cross-references: GB:006452; NID:9476131; PIDN:AAA19238.1; PID:9476132
R:Conliffe, P.G.; Brichard, V.; Van Pel, A.; Wolfel, T.; Schneider, J.; Traversari, L.
J. Exp. Med. 180, 35-42, 1994
A>Title: A new gene coding for a differentiation antigen recognized by autologous
A:Reference number: 138506; MUID:94275389; PMID:8006593
A:Accession: 138506
A>Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: mRNA
A:Residues: 1-118 <RES>
A:Cross-references: EMBL:U06654; NID:9517022; PIDN:AAA20389.1; PID:9517023
C:Genetics:
A:Gene: GDB:MLANA
A:Cross-references: GDB:358979
A:Map position: 17q21-17q24
C:Superfamily: human melanoma antigen MART-1

Query Match 100.0%; Score 649; DB 2; Length 118;
Best Local Similarity 100.0%; Pred. No. 3.5e-61;
Matches 118; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 MPREDAHRTGYPKKGHSHSTTAEAGIGITVITLGVLLIGCYCRRRGYALMDK 60
|||||
Db 1 MPREDAHRTGYPKKGHSHSTTAEAGIGITVITLGVLLIGCYCRRRGYALMDK 60
OY 61 SLHVTGTCALTRRCPOEGFDHRDSKVSIOENKCEPVNAPPAYEKLSEOSPPPYSP 118
|||||
Db 61 SLHVTGTCALTRRCPOEGFDHRDSKVSIOENKCEPVNAPPAYEKLSEOSPPPYSP 118

RESULT 2

T03455
ALR protein - human
C:Species: Homo sapiens (man)
C>Date: 24-Mar-1999 #sequence_revision 24-Mar-1999 #text_change 21-Jul-2000
R:Prasad, R.; Zhadanov, A.B.; Sedkov, Y.; Bullrich, F.; Druck, T.; Rallapalli, R.;
Oncogene 15, 549-560, 1997

A:Title: Structure and expression pattern of human ALR, a novel gene with strong homolog
A:Reference number: Z14954; MUID:97388474; PMID:9247308
A:Accession: T03455
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: mRNA
A:Residues: 1-4957 <PRA>
A:Cross-references: EMBL:AF010404; NID:g2358286; PIDN:AAC51735.1; PID:g2358287
C:Genetics:
A:Gene: ALR
A:Map position: 12
C:Superfamily: human ALR protein
C:Keywords: alternative splicing

Query Match 13.4%; Score 87; DB 2; Length 4957;
Best Local Similarity 31.2%; Pred. No. 3.2;
Matches 25; Conservative 7; Mismatches 30; Indels 18; Gaps 3;

OY 41 LITGWCRRRRNGRRLMDKSLHGTQCALTR---RCPOGPFDRDSKVSLOEKNCPEV 96
DB 939 LITGCRRCER-----WMHAGCESLFTEDVDNAHPDGF-----CVSCQPPYVAVPV 984

OY 97 VPNAPEYKLSAEOSSPPY 116
DB 985 APVAPPELVPMKVEPEPOY 1004

RESULT 3
ALR protein - human
C:Species: Homo sapiens (man)
C:Date: 24-Mar-1999 #sequence_revision 24-Mar-1999 #text_change 21-Jul-2000
C:Accession: T03454
R:Prasad, R.; Zhadanov, A.B.; Sedkov, Y.; Bullrich, F.; Druck, T.; Rallapalli, R.; Yano, Oncogene 15, 549-560, 1997
A:Title: Structure and expression pattern of human ALR, a novel gene with strong homolog
A:Reference number: Z14954; MUID:97388474; PMID:9247308
A:Accession: T03454
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: mRNA
A:Residues: 1-5262 <PRA>
A:Cross-references: EMBL:AF010403; NID:g2358284; PIDN:AAC51734.1; PID:g2358285
C:Genetics:
A:Gene: ALR
A:Map position: 12
C:Superfamily: human ALR protein
C:Keywords: alternative splicing

Query Match 13.4%; Score 87; DB 2; Length 5262;
Best Local Similarity 31.2%; Pred. No. 3.4;
Matches 25; Conservative 7; Mismatches 30; Indels 18; Gaps 3;

OY 41 LITGWCRRRRNGRRLMDKSLHGTQCALTR---RCPOGPFDRDSKVSLOEKNCPEV 96
DB 1244 LITGCRRCER-----WMHAGCESLFTEDVDNAHPDGF-----CVSCQPPYVAVPV 1289

OY 97 VPNAPEYKLSAEOSSPPY 116
DB 1290 APVAPPELVPMKVEPEPOY 1309

RESULT 4
B28967
T-cell surface glycoprotein CD2 precursor - mouse
N:Alternate names: CD2 antigen; T-lymphocyte antigen CD2; T11 protein
C:Species: Mus musculus (house mouse)
C:Date: 30-Jun-1989 #sequence_revision 03-Jun-1993 #text_change 23-Jul-1999
C:Accession: B28967; S01347; S02293
R:Diamond, D.J.; Clayton, L.K.; Sayre, P.H.; Reinherz, E.L.
Proc. Natl. Acad. Sci. U.S.A. 85, 1615-1619, 1988
A:Title: Exon-intron organization and sequence comparison of human and murine T11 (CD2)
A:Reference number: A28967; MUID:88144486; PMID:2894031
A:Accession: B28967
A:Molecule type: mRNA

A:Residues: 1-344 <CD1A>
A:Cross-references: GB:M19807; NID:g192479; PIDN:AAA37393.1; PID:g387122; GB:J03622
A:Note: the authors translated the codon TAT for residue 99 as Thr
R:Clayton, L.K.; Sayre, P.H.; Novotny, J.; Reinherz, E.L.
Eur. J. Immunol. 17, 1367-1370, 1987
A:Title: Murine and human T11 (CD2) cDNA sequences suggest a common signal transduc
A:Reference number: S01347; MUID:88004738; PMID:2820751
A:Accession: S01347
A:Molecule type: mRNA
A:Residues: 1-127, 'M', 129-174, 'N', 176-190, 'NM', 193-344 <CD1A>
A:Cross-references: EMBL:X06143; NID:g54223; PIDN:CAA29500.1; PID:g54224
R:Sewell, W.A.; Brown, M.H.; Owen, M.J.; Fluk, P.J.; Kozak, C.A.; Crumpton, M.J.
Eur. J. Immunol. 17, 1015-1020, 1987
A:Title: The murine homologue of the T lymphocyte CD2 antigen: molecular cloning, c
A:Reference number: S02293; MUID:87276135; PMID:2440689
A:Accession: S02293
A:Status: not compared with conceptual translation
A:Molecule type: mRNA
A:Residues: 1-127, 'M', 129-174, 'N', 176-191, 'M', 193-344 <SEM>
A:Cross-references: EMBL:Y00023; NID:g50346; PIDN:CAA68258.1; PID:g50347
C:Genetics:
A:Map position: 3
C:Superfamily: T-cell surface glycoprotein CD2
C:Keywords: glycoprotein; surface antigen; T-cell; transmembrane protein
F:1-22/Domain: signal sequence #status predicted <SIG>
F:23-344/Product: T-cell surface glycoprotein CD2 #status predicted <MAT>
F:23-203/Domain: extracellular #status predicted <EXT>
F:204-228/Domain: transmembrane #status predicted <TM>
F:229-344/Domain: intracellular #status predicted <INT>

Query Match 12.4%; Score 80.5; DB 2; Length 344;
Best Local Similarity 27.4%; Pred. No. 0.95;
Matches 31; Conservative 15; Mismatches 46; Indels 21; Gaps 5;

OY 3 REDAFHYGPKKGHSYTTAEAGIGILVYIGVLLITGWCRRRRNGRRLMDKSL 62
DB 188 KESTEVNCPKESGF-YVTVGVGAG-GLLVTL-VALFIPC-ICKRRRRRRRDEEL 243

OY 63 HVTQCALTRRCPOGPFDRDSKVSLOEKNCPEVVPNAPEYKLSAEOSSPP 115
DB 244 EI-----KASRTSTVERGPPHSTPAALAAONSVALQAPPP 279

RESULT 5
149585
CD2 antigen protein precursor - mouse
C:Species: Mus musculus (house mouse)
C:Date: 02-Jul-1996 #sequence_revision 02-Jul-1996 #text_change 23-Jul-1999
C:Accession: I49585
R:Yagita, H.; Okumura, K.; Nakauchi, H.
J. Immunol. 140, 1321-1326, 1988
A:Title: Molecular cloning of the murine homologue of CD2: Homology of the molecule
A:Reference number: I49585; MUID:88140313; PMID:3257775
A:Accession: I49585
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: mRNA
A:Residues: 1-344 <RES>
A:Cross-references: GB:M18934; NID:g192486; PIDN:AAA37397.1; PID:g309158
C:Superfamily: T-cell surface glycoprotein CD2

Query Match 12.4%; Score 80.5; DB 2; Length 344;
Best Local Similarity 27.4%; Pred. No. 0.95;
Matches 31; Conservative 15; Mismatches 46; Indels 21; Gaps 5;

OY 3 REDAFHYGPKKGHSYTTAEAGIGILVYIGVLLITGWCRRRRNGRRLMDKSL 62
DB 188 KESTEVNCPKESGF-YVTVGVGAG-GLLVTL-VALFIPC-ICKRRRRRRRDEEL 243

OY 63 HVTQCALTRRCPOGPFDRDSKVSLOEKNCPEVVPNAPEYKLSAEOSSPP 115
DB 244 EI-----KASRTSTVERGPPHSTPAALAAONSVALQAPPP 279

RESULT 6
T49773

related to actin-interacting protein AIP3 [Imported] - Neurospora crassa

N:Alternate names: protein B9J10.100

C:Species: Neurospora crassa

C>Date: 02-Jun-2000 #sequence_revision 02-Jun-2000 #text_change 02-Jun-2000

R:Schulte, U.; Aign, V.; Hohelsel, J.; Brandt, P.; Fartmann, B.; Holland, R.; Nyakatura, submitted to the Protein Sequence Database, May 2000

A:Reference number: 225022

A:Accession: T49773

A:Status: preliminary

A:Molecule type: DNA

A:Residues: 1-972 <SCH>

A:Cross-references: EMBL:AL356324; GSPDB:GN00116; NCSP:B9J10.100

A:Experimental source: BAC clone B9J10; strain OR74A

C:Genetics:

A:Gene: NCSP:B9J10.100

A:Map position: 6

A:Introns: 29/3; 161/1; 329/1

Query Match

Best Local Similarity

Matches

26; Conservative

10; Mismatches

31; Indels

18; Gaps

4;

12.2%; Score 79; DB 2; Length 972;

30.6%; Pred. No. 4.1;

10; Mismatches

31; Indels

18; Gaps

4;

12.2%; Score 79; DB 2; Length 972;

30.6%; Pred. No. 4.1;

10; Mismatches

31; Indels

18; Gaps

4;

12.2%; Score 79; DB 2; Length 972;

30.6%; Pred. No. 4.1;

10; Mismatches

31; Indels

18; Gaps

4;

12.2%; Score 79; DB 2; Length 972;

30.6%; Pred. No. 4.1;

10; Mismatches

31; Indels

18; Gaps

A:Gene: nsc

C:Superfamily: formate dehydrogenase

C:Keywords: 4Fe-4S; chromoprotein; iron-sulfur protein; metalloprotein; molybdenum

F;26/29/33,63/Binding site: 4Fe-4S cluster (Cys) (covalent) #status predicted

F;154/Binding site: molybdopterin (Cys) (covalent) #status predicted

Query Match

Best Local Similarity

Matches

34; Conservative

8; Mismatches

37; Indels

26; Gaps

6;

11.9%; Score 77; DB 2; Length 710;

32.4%; Pred. No. 4.8;

8; Mismatches

37; Indels

26; Gaps

6;

11.9%; Score 77; DB 2; Length 710;

32.4%; Pred. No. 4.8;

8; Mismatches

37; Indels

26; Gaps

6;

11.9%; Score 77; DB 2; Length 710;

32.4%; Pred. No. 4.8;

8; Mismatches

37; Indels

26; Gaps

6;

11.9%; Score 77; DB 2; Length 710;

32.4%; Pred. No. 4.8;

8; Mismatches

37; Indels

26; Gaps

6;

11.9%; Score 77; DB 2; Length 710;

32.4%; Pred. No. 4.8;

8; Mismatches

37; Indels

26; Gaps

6;

11.9%; Score 77; DB 2; Length 710;

32.4%; Pred. No. 4.8;

8; Mismatches

37; Indels

26; Gaps

A:Gene: nsc

C:Superfamily: formate dehydrogenase

C:Keywords: 4Fe-4S; chromoprotein; iron-sulfur protein; metalloprotein; molybdenum

F;26/29/33,63/Binding site: 4Fe-4S cluster (Cys) (covalent) #status predicted

F;154/Binding site: molybdopterin (Cys) (covalent) #status predicted

Query Match

Best Local Similarity

Matches

34; Conservative

8; Mismatches

37; Indels

26; Gaps

6;

11.9%; Score 77; DB 2; Length 710;

32.4%; Pred. No. 4.8;

8; Mismatches

37; Indels

26; Gaps

6;

11.9%; Score 77; DB 2; Length 710;

32.4%; Pred. No. 4.8;

8; Mismatches

37; Indels

26; Gaps

6;

11.9%; Score 77; DB 2; Length 710;

32.4%; Pred. No. 4.8;

8; Mismatches

37; Indels

26; Gaps

6;

11.9%; Score 77; DB 2; Length 710;

32.4%; Pred. No. 4.8;

8; Mismatches

37; Indels

26; Gaps

6;

11.9%; Score 77; DB 2; Length 710;

32.4%; Pred. No. 4.8;

8; Mismatches

37; Indels

26; Gaps

6;

11.9%; Score 77; DB 2; Length 710;

32.4%; Pred. No. 4.8;

8; Mismatches

37; Indels

26; Gaps

A:Gene: nsc

C:Superfamily: formate dehydrogenase

C:Keywords: 4Fe-4S; chromoprotein; iron-sulfur protein; metalloprotein; molybdenum

F;26/29/33,63/Binding site: 4Fe-4S cluster (Cys) (covalent) #status predicted

F;154/Binding site: molybdopterin (Cys) (covalent) #status predicted

Query Match

Best Local Similarity

Matches

34; Conservative

8; Mismatches

37; Indels

26; Gaps

6;

11.9%; Score 77; DB 2; Length 710;

32.4%; Pred. No. 4.8;

8; Mismatches

37; Indels

26; Gaps

6;

11.9%; Score 77; DB 2; Length 710;

32.4%; Pred. No. 4.8;

8; Mismatches

37; Indels

26; Gaps

6;

11.9%; Score 77; DB 2; Length 710;

32.4%; Pred. No. 4.8;

8; Mismatches

37; Indels

26; Gaps

6;

11.9%; Score 77; DB 2; Length 710;

32.4%; Pred. No. 4.8;

8; Mismatches

37; Indels

26; Gaps

6;

11.9%; Score 77; DB 2; Length 710;

32.4%; Pred. No. 4.8;

8; Mismatches

37; Indels

26; Gaps

6;

11.9%; Score 77; DB 2; Length 710;

32.4%; Pred. No. 4.8;

8; Mismatches

37; Indels

26; Gaps

A:Gene: nsc

C:Superfamily: formate dehydrogenase

C:Keywords: 4Fe-4S; chromoprotein; iron-sulfur protein; metalloprotein; molybdenum

F;26/29/33,63/Binding site: 4Fe-4S cluster (Cys) (covalent) #status predicted

F;154/Binding site: molybdopterin (Cys) (covalent) #status predicted

Query Match

Best Local Similarity

Matches

34; Conservative

8; Mismatches

37; Indels

26; Gaps

6;

11.9%; Score 77; DB 2; Length 710;

32.4%; Pred. No. 4.8;

8; Mismatches

37; Indels

26; Gaps

6;

11.9%; Score 77; DB 2; Length 710;

32.4%; Pred. No. 4.8;

8; Mismatches

37; Indels

26; Gaps

6;

11.9%; Score 77; DB 2; Length 710;

32.4%; Pred. No. 4.8;

8; Mismatches

37; Indels

26; Gaps

6;

11.9%; Score 77; DB 2; Length 710;

32.4%; Pred. No. 4.8;

8; Mismatches

37; Indels

26; Gaps

6;

11.9%; Score 77; DB 2; Length 710;

32.4%; Pred. No. 4.8;

8; Mismatches

37; Indels

26; Gaps

6;

11.9%; Score 77; DB 2; Length 710;

32.4%; Pred. No. 4.8;

8; Mismatches

37; Indels

26; Gaps

A:Gene: nsc

C:Superfamily: formate dehydrogenase

C:Keywords: 4Fe-4S; chromoprotein; iron-sulfur protein; metalloprotein; molybdenum

F;26/29/33,63/Binding site: 4Fe-4S cluster (Cys) (covalent) #status predicted

F;154/Binding site: molybdopterin (Cys) (covalent) #status predicted

Query Match

Best Local Similarity

Matches

34; Conservative

8; Mismatches

37; Indels

26; Gaps

6;

11.9%; Score 77; DB 2; Length 710;

32.4%; Pred. No. 4.8;

8; Mismatches

37; Indels

26; Gaps

6;

11.9%; Score 77; DB 2; Length 710;

32.4%; Pred. No. 4.8;

8; Mismatches

37; Indels

26; Gaps

6;

11.9%; Score 77; DB 2; Length 710;

32.4%; Pred. No. 4.8;

8; Mismatches

```

Query Match      37; Conservative   10; Mismatches    52; Indels     59; Gaps      7;
Best Local Similarity 25.5%; Pred. No. 31;

Db              37; Conservative   10; Mismatches    52; Indels     59; Gaps      7;
C:Species: Homo sapiens (man)
C:Date: 22-Jan-1999 #sequence_revision 22-Jan-1999 #text_change 01-Dec-2000
C:Accession: T00040
R:Yoshida, K.; Yoshimoto-Nakagawa, K.; Seki, N.; Sasaki, M.; Sugano, S.
Genomics 49, 458-461, 1998
A:title: Cloning, expression analysis, and chromosomal localization of BH-protocadherin
A:Reference number: Z14074; MID:98277460; PMID:9615233
A:Accession: T00041
A>Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: mRNA
A:Residues: 1-1072 <TOS>
A:Cross-references: EMBL:AB006756; NID:g2979419; PIDN:BAA25195.1; PID:g2979420
A:Experimental source: clone BH-Pcdh-b
A:Accession: T00040
A>Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: mRNA
A:Residues: 1-1058,'M',1060,'LH',1063,'Y',1065,'TVFG'<TO2>
A:Cross-references: EMBL:AB006755; NID:g2979417; PIDN:BAA25194.1; PID:g2979418
A:Experimental source: clone BH-Pcdh-a
C:Genetics:
A:Map position: 4p15

Query Match      10.9%; Score 71; DB 2; Length 1072;
Best Local Similarity 25.5%; Pred. No. 31;

Db              10.9%; Score 71; DB 2; Length 1072;
C:Species: Rhodospirillum rubrum
C:Date: 18-Aug-2000 #sequence_revision 18-Aug-2000 #text_change 18-Aug-2000
C:Accession: T51314
R:Kerby, R.L.
submitted to the EMBL Data Library, July 1996
A:Reference number: Z25372
A:Accession: T51314
A>Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-1265 <KER>
A:Cross-references: EMBL:U65510; PIDN:AAC45116.1
A:Experimental source: strain URI
C:Genetics:
A:Gene: cooM

Query Match      11.0%; Score 71.5; DB 2; Length 1265;
Best Local Similarity 30.9%; Pred. No. 33;
Matches 25; Conservativity 8; Mismatches 33; Indels 15; Gaps 3;

Db              11.0%; Score 71.5; DB 2; Length 1265;
C:Species: Homo sapiens (man)
C:Date: 22-Jan-1999 #sequence_revision 22-Jan-1999 #text_change 01-Dec-2000
C:Accession: T00040
R:Yoshida, K.; Yoshimoto-Nakagawa, K.; Seki, N.; Sasaki, M.; Sugano, S.
Genomics 49, 458-461, 1998
A:title: Cloning, expression analysis, and chromosomal localization of BH-protocadherin
A:Reference number: Z14074; MID:98277460; PMID:9615233
A:Accession: T00041
A>Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: mRNA
A:Residues: 1-1072 <TOS>
A:Cross-references: EMBL:AB006756; NID:g2979419; PIDN:BAA25195.1; PID:g2979420
A:Experimental source: clone BH-Pcdh-b
A:Accession: T00040
A>Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: mRNA
A:Residues: 1-1058,'M',1060,'LH',1063,'Y',1065,'TVFG'<TO2>
A:Cross-references: EMBL:AB006755; NID:g2979417; PIDN:BAA25194.1; PID:g2979418
A:Experimental source: clone BH-Pcdh-a
C:Genetics:
A:Map position: 4p15

Query Match      10.9%; Score 71; DB 2; Length 1072;
Best Local Similarity 25.5%; Pred. No. 31;

Db              10.9%; Score 71; DB 2; Length 1072;
C:Species: Rhodospirillum rubrum
C:Date: 18-Aug-2000 #sequence_revision 18-Aug-2000 #text_change 18-Aug-2000
C:Accession: T51314
R:Kerby, R.L.
submitted to the EMBL Data Library, July 1996
A:Reference number: Z25372
A:Accession: T51314
A>Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-1265 <KER>
A:Cross-references: EMBL:U65510; PIDN:AAC45116.1
A:Experimental source: strain URI
C:Genetics:
A:Gene: cooM

Query Match      11.0%; Score 71.5; DB 2; Length 1265;
Best Local Similarity 30.9%; Pred. No. 33;
Matches 25; Conservativity 8; Mismatches 33; Indels 15; Gaps 3;

Db              11.0%; Score 71.5; DB 2; Length 1265;
C:Species: Homo sapiens (man)
C:Date: 22-Jan-1999 #sequence_revision 22-Jan-1999 #text_change 01-Dec-2000
C:Accession: T00040
R:Yoshida, K.; Yoshimoto-Nakagawa, K.; Seki, N.; Sasaki, M.; Sugano, S.
Genomics 49, 458-461, 1998
A:title: Cloning, expression analysis, and chromosomal localization of BH-protocadherin
A:Reference number: Z14074; MID:98277460; PMID:9615233
A:Accession: T00041
A>Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: mRNA
A:Residues: 1-1072 <TOS>
A:Cross-references: EMBL:AB006756; NID:g2979419; PIDN:BAA25195.1; PID:g2979420
A:Experimental source: clone BH-Pcdh-b
A:Accession: T00040
A>Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: mRNA
A:Residues: 1-1058,'M',1060,'LH',1063,'Y',1065,'TVFG'<TO2>
A:Cross-references: EMBL:AB006755; NID:g2979417; PIDN:BAA25194.1; PID:g2979418
A:Experimental source: clone BH-Pcdh-a
C:Genetics:
A:Map position: 4p15

```

[illegible]

```

Db      113 TGSAMSTERENLRVLDELRLHAPQ---RTRLPPG---DSRL 150
          |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |

```

RESULT 14

688640
protein F52C12.3 [imported] - *Caenorhabditis elegans*
C:Species: *Caenorhabditis elegans*
C:Date: 10-May-2001 #sequence_revision 10-May-2001 #text_change 10-May-2001
C:Accession: G68640
R:Anonymous, The C. elegans Sequencing Consortium.
Science 282, 2012-2018, 1998
A:Title: Genome sequence of the nematode C. elegans: a platform for investigating biology
A:Reference number: A75000; MID:J99069613; PMID:9851916
A:Note: see websites genome.wustl.edu/gsc/C.elegans/ and www.sanger.ac.uk/Projects/C.elegans/
A:Accession: G68640.
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-335 <STO>
A:Cross-references: GB:chr_IV; PIDN:AA68961.1; PID:g3800953; GSPDB:GN00022; CESP:F52C12.3
C:Genetics:
A:Gene: F52C12.3
A:Map position: 4

RESULT 15

H84512
Hypothetical protein At2g14000 [imported] - Arabidopsis thaliana
C:Species: Arabidopsis thaliana (mouse-ear cress)
C:Date: 02-Feb-2001 #sequence_revision 02-Feb-2001 #text_change 02-Feb-2001
C:Accession: H84512
R:Lin, X.; Kaul, S.; Rounsley, S.D.; Shea, T.P.; Benito, M.I.; Town, C.D.; Fujii, C.Y.;
M.L.; Koo, H.; Moffat, K.S.; Cronin, L.A.; Shen, M.; Vankken, S.E.; Umayam, L.; Tallon, L.;
Euss, D.; Nierman, W.C.; White, O.; Eisen, J.A.; Salzberg, S.L.; Fraser, C.M.; Venter,
Nature 402, 761-769, 1999
A:Title: Sequence and analysis of chromosome 2 of the plant Arabidopsis thaliana.
A:Reference number: A84420; MUID:20083487; PMID:10617197
A:Accession: H84512
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-346 <STO>
A:Cross-references: GB:AE002093; NID:g4388823; PIDN:ADD19778.1; GSPDB:GN00139
C:Genetics:
A:Gene: At2g14000
A:Map position: 2

Db 331 GASTSQPPPHS 342

Search completed: October 7, 2003, 18:49:41
Job time : 21 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: October 7, 2003, 18:39:49 : Search time 41 Seconds
(Without alignments) 456.823 Million cell updates/sec

Title: US-09-898-860-2
Perfect score: 649
Sequence: 1 MPREDAHFTYGYRKQGHSH.....NAPPAVEKLSAQSPPPYSP 118

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 1107863 seqs, 158726573 residues

Total number of hits satisfying chosen parameters: 1107863

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%

Listing first 45 summaries

Database :
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2:	/SIDSL/gcgdata/geneseq/geneseq-emb1/AA1981.DAT:*
3:	/SIDSL/gcgdata/geneseq/geneseq-emb1/AA1982.DAT:*
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6:	/SIDSL/gcgdata/geneseq/geneseq-emb1/AA1985.DAT:*
7:	/SIDSL/gcgdata/geneseq/geneseq-emb1/AA1986.DAT:*
8:	/SIDSL/gcgdata/geneseq/geneseq-emb1/AA1987.DAT:*
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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	649	100.0	118	15	AAR63158 Tumour rejection a
2	649	100.0	118	16	AAR84212 MART-1 melanoma an
3	649	100.0	118	20	AAV42634 Human melanoma ant
4	649	100.0	118	20	AAAR83134 Human tumour rejec
5	649	100.0	118	22	AAU28888 MART-1 antigen. H
6	649	100.0	118	22	AA886042 Human MART1/Melana
7	649	100.0	118	23	AAU77793 Human melanoma ant
8	649	100.0	118	23	AAU84804 Human MART consens
9	649	100.0	118	23	AAU11541 Human Melanoma tum

10	649	100.0	119	20	AAV42633 Human melanoma ant
11	649	100.0	140	24	ABG76339 Recombinant mouse
12	649	100.0	496	24	ABG76343 Mouse recombinant
13	645	99.4	118	23	AAU77810 Human MART-1 prote
14	643	99.1	118	23	AAU77809 Human MART-1 prote
15	642	98.9	118	20	AAV31980 Human MART1 melano
16	641	98.8	118	23	AAU77806 Human MART-1 prote
17	641	98.8	118	23	AAU77808 Human MART-1 prote
18	641	98.8	118	23	AAU77811 Human MART-1 prote
19	640	98.6	118	23	AAU77807 Human MART-1 prote
20	599	92.3	118	23	AAU77815 Human MART-1 prote
21	598	92.1	118	23	AAU77813 Human MART-1 prote
22	598	92.1	118	23	AAU77814 Human MART-1 prote
23	597	92.0	118	23	AAU77812 Human MART-1 prote
24	596	91.8	118	23	AAU77816 Human MART-1 prote
25	562	86.6	104	23	AAU98926 Human melanoma ant
26	532	82.0	98	23	ABG70354 Novel human thromb
27	427.5	65.9	113	20	AAV31979 Mouse MART1 melano
28	427.5	65.9	114	20	AAV42632 Murine melanoma an
29	323	40.8	94	23	ABG70355 Novel human thromb
30	262.5	40.4	5546	23	AAU85008 Human MART segment
31	209	32.2	40	22	AAU86163 Human MART segment
32	173	26.7	30	23	AAU84868 Human MART segment
33	169	26.0	30	23	AAU84870 Human MART segment
34	167	25.7	30	23	AAU84871 Human MART segment
35	163	25.1	30	23	AAU84869 Human MART segment
36	161	24.8	30	23	AAU84867 Human MART segment
37	158	24.3	30	23	AAU84865 Human MART segment
38	156	24.0	30	22	AAU84864 MART 1 peptide #2.
39	148	22.8	30	23	AAU84866 Human MART segment
40	121	18.6	23	23	ABG79128 Human MART-1 class
41	108	16.6	21	18	AAU00903 Human melanoma tum
42	108	16.6	21	23	AAE20402 Human melanoma tum
43	98	15.1	22	23	AAU11546 Human control pept
44	82	12.6	17	23	AAU84872 Human MART segment
45	79	12.2	1114	22	ABU66628 Drosophila melanog

ALIGNMENTS

RESULT 1
ID AAR63158 standard; Protein: 118 AA.
XX
AC AAR63158;
XX
DT 25-MAR-2003 (updated)
DT 26-MAR-1995 (first entry)
XX
DE Tumour rejection antigen precursor.
XX
KW Tumour rejection antigen; precursor; HLA-A2 molecule; tyrosinase;
KW Isolation; melanoma; cell line; LB-39-MEL; diagnosis; vaccine;
KW therapy.
XX
OS Homo sapiens.
XX
PN WO9421126-A1.
XX
PD 29-SEP-1994.
XX
PF 09-MAR-1994; 94WO-US02487.
XX
PR 18-MAR-1993; 93US-0032978.
XX
PA (LUDWIG-) LUDWIG INST CANCER RES.
XX Boon-Fallleur T, Brichard V, De Plaen E, Traversari C;
PI Van Pel A, Wolfel T;
XX WPI: 1994-316544/39.
XX N-PSDB: AAQ76370.
DR

XX Nucleic acid coding for a tumour rejection antigen precursor - is
 PT used for developing prods. for diagnosis or treatment of expression
 PT related disorders, partic. melanoma
 XX
 PS Claim 5; Page 14; 26pp; English.

CC This sequence represents the tumour rejection antigen precursor which is
 CC processed to a tumour rejection antigen presented by HLA-A2 molecules.
 CC The tumour rejection antigen is not related to tyrosinase. The CDNA
 CC encoding this sequence was isolated from the melanoma cell line,
 CC LB-39-MEL. The tumour rejection antigen may be used for diagnosis or
 CC in vaccines or for therapy of disorders characterised by the expression
 CC of the tumour rejection antigen precursor, particularly melanoma.
 CC (Updated on 25-MAR-2003 to correct PN field.)
 CC
 XX

SQ Sequence 118 AA;

Query Match 100.0%; Score 649; DB 15; Length 118;
 Best Local Similarity 100.0%; Pred. No. 2,1e-65;
 Matches 118; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MPRDAHFHYGYPKKGHSYTTAEAGIGILTVILGVLILGICWYCRRRNGYRALMDK 60
 DB 1 MPRDAHFHYGYPKKGHSYTTAEAGIGILTVILGVLILGICWYCRRRNGYRALMDK 60

QY 61 SLHVGTCALTRRCPOEGFDHRDHSKYSIQEKNCPEVVPNAPPAYEKLSAQSPPPYSP 118
 DB 61 SLHVGTCALTRRCPOEGFDHRDHSKYSIQEKNCPEVVPNAPPAYEKLSAQSPPPYSP 118

RESULT 2

AA84212
 ID AA84212 standard; Protein: 118 AA.

AC AA84212;

DT 20-APR-1996 (first entry)

DE MART-1 melanoma antigen.

XX MART-1; melanoma antigen recognised by T-cell; melanoma;
 KW metastatic melanoma; tumour-associated antigen; immunogen;
 KW diagnosis; prognosis; prophylaxis; therapy; vaccine.

XX Mammalian.

OS

EH Key Location/Qualifiers

FT Region 27..47 /note="hydrophobic region"

XX W09529193-A2.

XX 02-NOV-1995.

XX 21-APR-1995; 95WO-US05063.

XX 05-APR-1995; 95US-0417174.

XX 22-APR-1994; 94US-0231365.

XX (USSH) US SEC DEPT HEALTH.

XX Kawakami Y, Rosenberg SA;

XX WPI; 1995-382963/49.

XX N-PSDB; AAT02714.

XX DNA encoding melanoma antigens recognised by T-lymphocytes - also
 PT vectors, host cells and antibodies, used to detect, treat and
 PT immunise animal against melanoma.

XX Claim 11; Page 117; 184pp; English.

CC The melanoma antigen (MART-1) is produced by recombinant DNA
 CC methods, i.e. preferably using a baculovirus vector for expression
 CC in insect cell cultures. MART-1 protein is a source of immunogenic
 CC peptides (see AAR84196 for peptide M9-2) which are optionally modified
 CC (see AAR84783-R84800) and used in medicaments for the treatment or
 CC prevention (by immunization) of melanoma. Antibodies against MART-1
 CC and its immunogenic peptides may be used in the detection and
 CC isolation of MART-1 from a sample, the detection of which is
 CC indicative of a disease state (melanoma or metastatic melanoma).
 CC
 XX

SQ Sequence 118 AA;

Query Match 100.0%; Score 649; DB 16; Length 118;
 Best Local Similarity 100.0%; Pred. No. 2,1e-65;
 Matches 118; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MPRDAHFHYGYPKKGHSYTTAEAGIGILTVILGVLILGICWYCRRRNGYRALMDK 60
 DB 1 MPRDAHFHYGYPKKGHSYTTAEAGIGILTVILGVLILGICWYCRRRNGYRALMDK 60

QY 61 SLHVGTCALTRRCPOEGFDHRDHSKYSIQEKNCPEVVPNAPPAYEKLSAQSPPPYSP 118
 DB 61 SLHVGTCALTRRCPOEGFDHRDHSKYSIQEKNCPEVVPNAPPAYEKLSAQSPPPYSP 118

RESULT 3

AA42634
 ID AA42634 standard; Protein: 118 AA.

AC AA42634;

DT 10-JAN-2000 (first entry)

DE Human melanoma antigen hMART1.

XX Immune response; self-antigen; immune effector cell; cancer; melanoma;
 KW mouse; melanoma antigen; MART1.

XX Homo sapiens.

XX W09946988-A1.

XX 23-SEP-1999.

XX 19-MAR-1999; 99WO-US06034.

XX 20-MAR-1998; 98US-0078890.

XX (GENZ) GENZYME CORP.

XX Nicolette CA;

XX WPI; 1999-580277/49.

XX N-PSDB; AA207987.

XX Method of inducing an immune reaction to a self-antigen by
 PT administering the antigen, especially useful for treating cancer or
 PT melanoma

XX Disclosure; Fig 3A-B; 70pp; English.

CC The invention provides a method of inducing a prophylactic immune
 CC response to a self-antigen in a subject. The method comprises
 CC administering the antigen or its derivative or administering educated
 CC immune effector cells able to recognize and lyse cells expressing the
 CC self-antigen or its derivative. The method is used to stimulate an immune
 CC response against a self-antigen especially one expressed in a cancer or
 CC melanoma. The present sequence represents the human melanoma antigen
 CC hMART1.

SQ Sequence 118 AA;

Query Match 100.0%; Score 649; DB 20; Length 118;

Best Local Similarity 100.0%; Pred. No. 2.1e-65;
Matches 118; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 MPREDAHFIYGYPKKGHSYTTAEAGIGILTVILGVLIIIGCWYCRRRNGYRALMDK 60
DB 1 MPREDAHFIYGYPKKGHSYTTAEAGIGILTVILGVLIIIGCWYCRRRNGYRALMDK 60
OY 61 SLHWGTQCALTRRCPOEGFDHRDHSKVSIOEKNCPEVVPNAPPAYEKLISAEGSPPPYSP 118
DB 61 SLHWGTQCALTRRCPOEGFDHRDHSKVSIOEKNCPEVVPNAPPAYEKLISAEGSPPPYSP 118

RESULT 4

AAW83134 ID AAW83134 standard; Protein; 118 AA.

AC AAW83134;

DT 04-FEB-1999 (first entry)

DE Human tumour rejection antigen precursor.

KW Human; tumour rejection antigen precursor; human leukocyte antigen;

KW TRAP; HLA; cancer; melanoma.

OS Homo sapiens.

EH Key Location/Qualifiers

FT Misc-difference 2 /note= "encoded by CGA"

FT Misc-difference 17 /note= "encoded by GNC"

FT US837476-A.

PN 17-NOV-1998.

PE 16-JAN-1998; 98US-0007966.

PR 03-MAR-1995; 95US-0398409.

PR 16-JAN-1998; 98US-0007966.

XX (LUDWIG) LUDWIG INST CANCER RES.

PI Boon-Falleur T, Brichard V, De Plaen E, Traversari C;

DR WPI; 1999-043967/04.

DR N-PSDB; AAV70150.

PT Use of a tumour rejection antigen precursor - as a marker for
diagnosing a disorder characterised by expression of a tumour
rejection antigen precursor which is not tyrosinase

PS Claim 1; Column 7-9; 11pp; English.

XX A method has been developed for the diagnosis of a disorder which is
characterised by the expression of a tumour rejection antigen precursor
(TRAP) which is not tyrosinase, and which is processed to a TRAP which
forms a complex with an HLA-A2 molecule. The present sequence represents
the TRAP for use in the present invention. The method comprises
contacting a sample from a subject with an agent specific for the
complex and determining the interaction between the complex and the
agent as a determination of the disorder. TRAP can be used for the
diagnosis and treatment of disorders characterised by the expression
of the TRAP molecules such as cancers, particularly melanoma.

SO Sequence 118 AA;

Query Match 100.0%; Score 649; DB 20; Length 118;
Best Local Similarity 100.0%; Pred. No. 2.1e-65;
Matches 118; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 MPREDAHFIYGYPKKGHSYTTAEAGIGILTVILGVLIIIGCWYCRRRNGYRALMDK 60
DB 1 MPREDAHFIYGYPKKGHSYTTAEAGIGILTVILGVLIIIGCWYCRRRNGYRALMDK 60
OY 61 SLHWGTQCALTRRCPOEGFDHRDHSKVSIOEKNCPEVVPNAPPAYEKLISAEGSPPPYSP 118
DB 61 SLHWGTQCALTRRCPOEGFDHRDHSKVSIOEKNCPEVVPNAPPAYEKLISAEGSPPPYSP 118

RESULT 5

AAU28888 ID AAU28888 standard; Protein; 118 AA.

AC AAU28888;

DT 18-DEC-2001 (first entry)

DE MART-1 antigen.

KW Human; MART-1; immunogenic; melanoma antigen recognised by T lymphocyte;

KW diagnostic; therapeutic; vaccine; melanoma; in vivo tumour recognition;

OS Homo sapiens.

PN US6270778-B1.

PD 07-AUG-2001.

PE 12-MAR-1999; 99US-0267439.

PR 05-MAY-1998; 98US-0073138.

PR 22-APR-1994; 94US-0231565.

PR 05-APR-1995; 95US-0417174.

XX (USSH) US DEPT HEALTH & HUMAN SERVICES.

PI Kawakami Y, Rosenberg SA;

DR WPI; 2001-595403/67.

DR N-PSDB; AAS45524.

PT Immunogenic peptide useful in vaccines comprises specific amino acids
of new melanoma antigen recognised by T lymphocytes

PS Claim 2; Figure 1; 73pp; English.

XX The invention relates to a novel immunogenic peptide comprising 5-20
contiguous amino acids of new melanoma antigen recognised by T
lymphocytes (MART-1). The peptide sequence contains at least one amino
acid modification of MART-1. The peptide is used in diagnostic and
therapeutic methods as an immunogen or vaccine to prevent or treat
melanoma, and for in vivo tumour recognition and rejection. AAU28888-
AAU29008 represent MART-1 peptide amino acid sequences, and related
sequences of the invention.

SO Sequence 118 AA;

Query Match 100.0%; Score 649; DB 22; Length 118;
Best Local Similarity 100.0%; Pred. No. 2.1e-65;
Matches 118; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 MPREDAHFIYGYPKKGHSYTTAEAGIGILTVILGVLIIIGCWYCRRRNGYRALMDK 60
DB 1 MPREDAHFIYGYPKKGHSYTTAEAGIGILTVILGVLIIIGCWYCRRRNGYRALMDK 60
OY 61 SLHWGTQCALTRRCPOEGFDHRDHSKVSIOEKNCPEVVPNAPPAYEKLISAEGSPPPYSP 118
DB 61 SLHWGTQCALTRRCPOEGFDHRDHSKVSIOEKNCPEVVPNAPPAYEKLISAEGSPPPYSP 118

RESULT 6

AAAB6042


```

ID   AAB86042 standard; Protein; 118 AA.
XX
AC   AAB86042;
XX
DT   16-JUL-2001 (first entry)
XX
DE   Human MART1/Melana protein.
XX
KW   Listeria: expression vector; tumor-associated antigen; Trp 1; Trp 2;
KW   Melana/MART-1; cytosolic; attenuated; immunotherapy; malignant melanoma;
KW   pigmented tumor; malignant schwannoma; vaccination; tyrosinase;
KW   antigen-presenting cell.
XX
OS   Homo sapiens.
XX
PN   WO200127295-A1.
XX
PD   19-APR-2001.
XX
PE   13-OCT-2000; 2000WO-DE03629.
XX
PR   14-OCT-1999; 99DE-1049594.
XX
PA   (DEKR-) DEUT KREBSFORSCHUNGSZENTRUM.
XX
PI   Schenendorf D, Paschen A, Chakraborty T, Dommann E;
XX
DR   WPI; 2001-282041/29.
XX
DR   N-PSDB; AAF86044.
XX
PT   Listeria expression vector for immunotherapy, particularly of malignant
PT   melanoma, comprises a DNA sequence encoding tumor-associated antigens -
XX
XX
PS   Disclosure; Fig 4; 41pp: German.
XX
CC   This invention describes a novel Listeria expression vector (A) for
CC   immunotherapy which comprises a promoter (P), functional in Listeria,
CC   operably linked to a DNA sequence (I) encoding one of the
CC   tumor-associated antigens (II) human tyrosinase, Trp 1 or 2, or
CC   Melana/MART-1. The products of the invention have cytostatic activity.
CC   Recombinant attenuated Listeria containing (A) are useful for
CC   immunotherapy (prophylactic, adjuvant or therapeutic), specifically of
CC   malignant melanoma (but also other pigmented tumors such as malignant
CC   schwannoma), particularly as a replacement for radiotherapy. Using
CC   attenuated Listeria as carrier for (A) provides a simple way of
CC   vaccination, since antigen-presenting cells acquire tumor-associated
CC   antigens by natural infection, eliminating the need for labor-intensive
CC   ex vivo modification of autologous cells. This sequence represents the
CC   human MART-1/Melana protein described in the method of the invention.
XX
SQ   Sequence 118 AA;
Query Match 100.0%; Score 649; DB 22; Length 118;
Best Local Similarity 100.0%; Pred. No. 2.1e-65;
Matches 118; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 MPREDAHFTYGYPKKGHSYTTAEAGIGILTVTLIGVLLIGWCYCRRRNGYRLMDK 60
DB 1 MPREDAHFTYGYPKKGHSYTTAEAGIGILTVTLIGVLLIGWCYCRRRNGYRLMDK 60
QY 61 SLHVGTCALTRRCPOEGFDHRDSKYSLOEKNCCEPVVNPAPAYEKLSAEOSSPPYSP 118
DB 61 SLHVGTCALTRRCPOEGFDHRDSKYSLOEKNCCEPVVNPAPAYEKLSAEOSSPPYSP 118
RESULT 7
ID AAV77793 standard; Protein; 118 AA.
XX
AC AAV77793;
XX
DT 05-JUN-2002 (first entry)
XX

```

```

DE   Human melanoma antigen recognised by T cells (MART-1).
XX
KW   MART-1; melanoma antigen recognised by T cells; human;
KW   anti-melanoma peptide; major histocompatibility complex; MHC;
KW   immunotherapy; cancer; vaccine; immunoregulatory.
XX
OS   Homo sapiens.
XX
PN   WO200212272-A2.
XX
PD   14-FEB-2002.
XX
PE   03-AUG-2001; 2001WO-US24328.
XX
PR   04-AUG-2000; 2000US-223641P.
XX
PR   13-DEC-2000; 2000US-255502P.
XX
PR   25-JAN-2001; 2001US-264432P.
XX
PR   26-MAR-2001; 2001US-279005P.
XX
PA   (GENZ ) GENZYME CORP.
XX
PI   Nicolette CA;
XX
DR   WPI; 2002-257459/30.
XX
DR   N-PSDB; ABR11763.
XX
PT   Novel anti-melanoma peptide compounds useful for inducing immune
PT   response in a subject, and in the preparation of medicaments for the
PT   treatment and diagnosis of cancer -
XX
XX
PS   Disclosure; Page 69-70; 79pp: English.
XX
CC   This invention relates to a novel anti-melanoma peptide compound
CC   comprising a peptide of the human melanoma antigen recognised by T cells
CC   (MART-1) protein. These compounds are designed to enhance binding to
CC   major histocompatibility (MHC) compounds and enhance immunoregulatory
CC   properties and induce an immune response. The invention also comprises
CC   the nucleotide sequences encoding the peptides of the invention. The
CC   compounds of the invention are useful for inducing an immune response in
CC   a subject, by delivering the compounds in the context of a major
CC   histocompatibility (MHC) molecule which presents the compound on the
CC   surface of an antigen presenting cell, or by delivering it as a
CC   polynucleotide that encodes the peptide. The invention also comprises
CC   antibodies that recognise and bind these compounds which are useful in
CC   immunotherapy. The compounds of the invention are useful for modulating
CC   an immune response to synthetic and naturally occurring compounds in a
CC   subject. The compounds may also be used as components of anti-cancer
CC   vaccines and to expand immune effector cells that are specific for
CC   cancers characterised by expression of the human melanoma antigen
CC   recognised by T cells, MART-1. The compounds of the invention are also
CC   useful for the detection and purification of antibodies and may be used
CC   for the preparation of medicaments for the diagnosis and treatment of
CC   diseases such as cancer. The compounds of the invention have enhanced
CC   binding to MHC molecules and enhanced immunoregulatory properties
CC   relative to their natural counterparts. The present sequence represents
CC   the human melanoma antigen recognised by T cells (MART-1) used to create
CC   the peptide compounds of the invention.
XX
SQ   Sequence 118 AA;
Query Match 100.0%; Score 649; DB 23; Length 118;
Best Local Similarity 100.0%; Pred. No. 2.1e-65;
Matches 118; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 MPREDAHFTYGYPKKGHSYTTAEAGIGILTVTLIGVLLIGWCYCRRRNGYRLMDK 60
DB 1 MPREDAHFTYGYPKKGHSYTTAEAGIGILTVTLIGVLLIGWCYCRRRNGYRLMDK 60
QY 61 SLHVGTCALTRRCPOEGFDHRDSKYSLOEKNCCEPVVNPAPAYEKLSAEOSSPPYSP 118
DB 61 SLHVGTCALTRRCPOEGFDHRDSKYSLOEKNCCEPVVNPAPAYEKLSAEOSSPPYSP 118

```

RESULT 8
 ID AA084804 standard; protein; 118 AA.
 XX AA084804;
 AC
 XX
 DT 08-MAY-2002 (first entry)
 XX
 DE Human MART consensus sequence.
 XX
 KW Savine; vaccine; cancer; viral infection; HIV; hepatitis C virus;
 KW viral infection; human immunodeficiency virus; melanoma;
 KW bacterial infection; Salmonella; Legionella; parasitic infection;
 KW Trypanosoma; Toxoplasma; Giardia.
 XX
 OS Homo sapiens.
 XX
 PN WO200190197-A1.
 XX
 PD 29-NOV-2001.
 XX
 PF 25-MAY-2001; 2001WO-AU00622.
 XX
 PR 26-MAY-2000; 2000AU-0007761.
 XX
 PA (AUSU) UNIV AUSTRALIAN NAT.
 XX
 PI Thomson SA, Ramsay IA;
 XX
 DR WPI: 2002-147575/19.
 XX
 PT New synthetic polypeptides having several different segments of at
 PT least one parent polypeptide linked together differently compared to
 PT the linkage in the parent polypeptide, for inducing immune response
 PT against a pathogen or cancer.
 XX
 PS Example 3; Fig 27; 364pp; English.
 XX
 CC The invention relates to a new synthetic polypeptide (1) comprising
 CC several different segments of at least one parent polypeptide linked
 CC together in a different relationship relative to their linkage in the
 CC parent polypeptide to impede, abrogate or otherwise alter at least one
 CC function associated with the parent polypeptide and for inducing an
 CC immune response against a pathogen or cancer. Also included are a
 CC synthetic polynucleotide encoding and a computer system for
 CC designing the synthetic polypeptides. The synthetic polypeptides and
 CC polynucleotides are referred to as a Savine. The synthetic polypeptide is
 CC useful for modulating immune responses preferably directed against a
 CC pathogen or a cancer, (e.g., cancers of the lung, breast, ovary, cervix,
 CC colon, head and neck, pancreas, prostate, stomach, bladder, kidney, bone
 CC liver, oesophagus, brain, testicle, uterus), as potentiating agents.
 CC Compositions comprising the polypeptide may be used in the treatment or
 CC prophylaxis against viral (such as infections caused by HIV (human
 CC immunodeficiency virus), hepatitis, influenza, Japanese encephalitis
 CC virus, Epstein-Barr virus and respiratory syncytial virus), bacterial
 CC (e.g., infections caused by Neisseria, Meningococcal, Haemophilus,
 CC Salmonella, Streptococcal, Legionella and Mycobacterium) or parasitic
 CC (e.g., infections caused by Plasmodium, Schistosoma, Leishmania,
 CC Trypanosoma, Toxoplasma and Giardia) infections. The present
 CC sequence is a consensus sequence for a parent protein used to design a
 CC savine of the invention.
 XX
 SQ Sequence 118 AA:
 Query Match 100.0%; Score 649; DB 23; Length 118;
 Best Local Similarity 100.0%; Pred. No. 2,1e-65;
 Matches 118; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 MPREDAHFTYGYPKKGSHSYTAAEAGIGLTVLIGVLLTGGCWCRRNRGRAMDK 60
 DB 1 MPREDAHFTYGYPKKGSHSYTAAEAGIGLTVLIGVLLTGGCWCRRNRGRAMDK 60
 QY 61 SLHVGTCALTRRCPOEGFDHRDSKVSLOENKCEPVVPNAPYAEKLSAQSPPTYP 118

DB 61 SLHVGTCALTRRCPOEGFDHRDSKVSLOENKCEPVVPNAPYAEKLSAQSPPTYP 118
 RESULT 9
 ID AA011541 standard; protein; 118 AA.
 XX AA011541;
 AC
 XX
 DT 12-MAR-2002 (first entry)
 XX
 DE Human Melanoma tumour associated antigen.
 XX
 KW Human; melanoma tumour associated antigen; human leukocyte antigen;
 KW major histocompatibility complex; MHC; HLA-A2.2; vaccine; cancer;
 KW HIV; human immunodeficiency virus infection; cytostatic; vitreous;
 KW housekeeping epitope; adoptive immunotherapy; neoplastic disease;
 KW viral disease; hepatitis virus; papilloma virus; tumour; leukaemia;
 KW lymphoma; breast cancer; prostate cancer; lung cancer; melan A;
 KW parasitic infection; Chlamydia; Trypanosoma; Toxoplasma.
 XX
 OS Homo sapiens.
 XX
 PN WO200182963-A2.
 XX
 PD 08-NOV-2001.
 XX
 PF 27-APR-2001; 2001WO-US13806.
 XX
 PR 28-APR-2000; 2000US-0560465.
 XX
 PR 28-APR-2000; 2000US-0561074.
 PR 28-APR-2000; 2000US-0561571.
 PR 28-APR-2000; 2000US-0561572.
 XX
 PA (CTLI-) CTL IMMUNOTHERAPIES CORP.
 XX
 PI Sismard JDL, Diamond DC, Lei X;
 XX
 DR WPI: 2002-066492/09.
 XX
 PT Novel vaccine useful for treating neoplastic and viral diseases,
 PT comprises a first housekeeping epitope derived from a first antigen
 PT associated with a first target cell.
 XX
 PS Example 21; Fig 15; 131pp; English.
 XX
 CC The invention relates to a vaccine comprising a first housekeeping
 CC epitope derived from a first antigen associated with a first target
 CC cell. Also included are an isolated T cell expressing a T cell receptor
 CC specific for a major histocompatibility complex (MHC)-peptide complex
 CC comprising a first housekeeping epitope which is derived from a first
 CC antigen associated with a first target cell, selecting an epitope
 CC (or peptide sequence) from a population of peptide fragments of an
 CC antigen associated with a target in a host, where the fragments have a
 CC known or predicted affinity for a MHC receptor peptide binding cleft of
 CC the host, where the epitope selected corresponds to a product of
 CC a nucleic acid construct comprising a first coding region, where the
 CC first coding region comprises a first sequence encoding at least a first
 CC polypeptide, where the first polypeptide comprises a first housekeeping
 CC epitope derived from a first antigen associated with a first target cell;
 CC The epitopes, peptides, vaccines and nucleic acids are useful in the
 CC manufacture of a medicament for use in adoptive immunotherapy and for
 CC prevention and treatment of neoplastic and viral diseases (e.g.,
 CC human immunodeficiency virus, HIV, infection, hepatitis virus and
 CC papilloma virus), cancers (e.g., tumours, leukaemia, lymphoma, breast
 CC cancer, prostate cancer and lung cancer), infection of cells by
 CC intracellular parasites (e.g., Chlamydia, Trypanosoma and
 CC Toxoplasma) and many other examples given in the specification.
 CC The invention permits the vaccine designer to ignore peptides that,
 CC despite predicted high binding affinity for MHC, will never be useful
 CC because they cannot be presented by target cells. The invention provides

CC a major advance in vaccine design, one that combines the power of antigen
 CC sequence analysis with the fundamental realities of immunology. The
 CC invention allows for the simple and effective selection of meaningful
 CC epitopes for creation of MHC class I or Class II vaccines using any
 CC polypeptide sequence corresponding to a desired target. The present
 CC sequence is an HLA-A2.1 (human leukocyte antigen) presenting target cell
 CC protein from which epitopes of the invention may be derived, Mela-A
 CC (melanoma tumour associated antigen).

XX SQ Sequence 118 AA;

Query Match 100.0%; Score 649; DB 23; Length 118;
 Best Local Similarity 100.0%; Pred. No. 2.1e-65;
 Matches 118; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 MPREDAHFIYGYPKKGHSHTTAEEAAGIGILTVILGVLILGICWYCRRRNGYRALMDK 60
 DB 1 MPREDAHFIYGYPKKGHSHTTAEEAAGIGILTVILGVLILGICWYCRRRNGYRALMDK 60
 OY 61 SLHVGTCALTRRCPOEGFDRHDSKVSLOEKNCPEVVPNAPPAYEKLSAEQSPPPYSP 118
 DB 61 SLHVGTCALTRRCPOEGFDRHDSKVSLOEKNCPEVVPNAPPAYEKLSAEQSPPPYSP 118

RESULT 10

AA42633 ID AA42633 standard; Protein: 119 AA.

XX AC AA42633;

XX DT 10-JAN-2000 (first entry)

XX DE Human melanoma antigen hMART1.

XX KM Immune response; self-antigen; immune effector cell; cancer; melanoma;
 XX mouse; melanoma antigen; MART1.

XX OS Homo sapiens.

XX FH Key Location/Qualifiers

XX FT Misc-difference 119 /note="unspecified"

XX PN WO9946988-A1.

XX PD 23-SEP-1999.

XX PF 19-MAR-1999; 99WO-US06034.

XX PR 20-MAR-1998; 98US-0078890.

XX PA (GENZ) GENZYME CORP.

XX PI Nicotlette CA;

XX DR WPI: 1999-580277/49.

XX N-PSDB; AAZ07986.

XX PT Method of inducing an immune reaction to a self-antigen by
 PT administering the antigen, especially useful for treating cancer or
 PT melanoma

XX PS Disclosure; Fig 2B; 70pp; English.

XX CC The invention provides a method of inducing a prophylactic immune

XX CC response to a self-antigen in a subject. The method comprises

XX CC administering the antigen or its derivative or administering educated

XX CC immune effector cells able to recognize and lyse cells expressing the

XX CC self-antigen or its derivative. The method is used to stimulate an immune
 CC response against a self-antigen especially one expressed in a cancer or
 CC melanoma. The present sequence represents the human melanoma antigen
 CC hMART1.

SQ Sequence 119 AA;

Query Match 100.0%; Score 649; DB 20; Length 119;
 Best Local Similarity 100.0%; Pred. No. 2.2e-65;
 Matches 118; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 MPREDAHFIYGYPKKGHSHTTAEEAAGIGILTVILGVLILGICWYCRRRNGYRALMDK 60
 DB 1 MPREDAHFIYGYPKKGHSHTTAEEAAGIGILTVILGVLILGICWYCRRRNGYRALMDK 60

OY 61 SLHVGTCALTRRCPOEGFDRHDSKVSLOEKNCPEVVPNAPPAYEKLSAEQSPPPYSP 118
 DB 61 SLHVGTCALTRRCPOEGFDRHDSKVSLOEKNCPEVVPNAPPAYEKLSAEQSPPPYSP 118

RESULT 11

ABG76339 ID ABG76339 standard; Protein: 140 AA.

XX AC ABG76339;

XX DT 10-MAY-2003 (first entry)

XX DE Recombinant mouse protein, Melana/MART1.

XX KM Mouse; protein targeting; exosome; lactadherin; C1 domain; C2 domain;
 XX membrane vesicle; mutant; mutein.

XX OS Mus sp.

XX OS Synthetic.

XX PN WO2003016522-A2.

XX PD 27-FEB-2003.

XX PF 14-AUG-2002; 2002WO-EP09108.

XX PR 17-AUG-2001; 2001US-313159P.

XX PR 26-DEC-2001; 2001US-343991P.

XX PA (ANOS) ANOSYS INC.

XX PI Delcayre A, Le Pecq J;

XX DR WPI: 2003-268331/26.

XX PT Targeting polypeptides to exosomes providing a chimeric genetic
 PT construct and introducing the construct into exosome-producing cells in
 PT vivo or ex vivo

XX PS Example 6; Page 85-86; 94pp; English.

XX CC The present invention relates to a method and compounds for targeting

XX CC polypeptides to exosomes. The method comprises providing a chimeric

XX CC genetic construct encoding the polypeptide fused to a targeting

XX CC polypeptide comprising lactadherin or its portion comprising

XX CC a functional C1 and/or C2 domain, and introducing the construct into

XX CC exosome-producing cells in vivo or ex vivo, to generate recombinant

XX CC vesicles. The method is useful for targeting proteins to membrane

XX CC vesicles, particularly exosomes, and is useful in experimental,
 CC research, therapeutic, prophylactic, and diagnostic areas. The
 CC present sequence represents a recombinant mouse protein.

XX SQ Sequence 140 AA;

Query Match 100.0%; Score 649; DB 24; Length 140;
 Best Local Similarity 100.0%; Pred. No. 2.6e-65;
 Matches 118; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 MPREDAHFIYGYPKKGHSHTTAEEAAGIGILTVILGVLILGICWYCRRRNGYRALMDK 60
 DB 1 MPREDAHFIYGYPKKGHSHTTAEEAAGIGILTVILGVLILGICWYCRRRNGYRALMDK 60

QY 61 SLHVGTCALTRRCPOEGFDHRDSKVSIOEKNCPEVVPNAPPAYEKLSAEO\$PPPYSP 118
 ID 61 SLHVGTCALTRRCPOEGFDHRDSKVSIOEKNCPEVVPNAPPAYEKLSAEO\$PPPYSP 118
 DB 61 SLHVGTCALTRRCPOEGFDHRDSKVSIOEKNCPEVVPNAPPAYEKLSAEO\$PPPYSP 118

RESULT 12

ABG76343
 ID ABG76343 standard; Protein; 496 AA.

AC ABG76343;

DT 10-MAY-2003 (first entry)

DE Mouse recombinant chimeric fusion protein, MART1/CCR7.

KW Mouse; protein targeting; exosome; lactadherin; C1 domain; C2 domain;
 KM membrane vesicle; mutant; mutein.

OS Mus sp.

PN WO2003016522-A2.

PD 27-FEB-2003.

PF 14-AUG-2002; 2002WO-EP09108.

PR 17-AUG-2001; 2001US-313159P.

PR 26-DEC-2001; 2001US-343991P.

PA (ANOS-) ANOSYS INC.

PI Delcayre A, Le Pecq J;

DR WPI; 2003-268331/26.

PT Targeting polypeptides to exosomes providing a chimeric genetic
 PT construct and introducing the construct into exosome-producing cells in
 PT vivo or ex vivo

PS Claim 28; Page 90-92; 94pp; English.

CC The present invention relates to a method and compounds for targeting
 CC polypeptides to exosomes. The method comprises providing a chimeric
 CC genetic construct encoding the polypeptide fused to a targeting
 CC polypeptide comprising lactadherin or its portion comprising
 CC a functional C1 and/or C2 domain, and introducing the construct into
 CC exosome-producing cells in vivo or ex vivo, to generate recombinant
 CC vesicles. The method is useful for targeting proteins to membrane
 CC vesicles, particularly exosomes, and is useful in experimental,
 CC research, therapeutic, prophylactic, and diagnostic areas. The
 CC present sequence represents a mouse recombinant chimeric fusion
 CC protein.

SO Sequence 496 AA;

Query Match 100.0%; Score 649; DB 24; Length 496;
 Best Local Similarity 100.0%; Pred. No. 1.3e-64;

Matches 118; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 MPREDAFITGYGPKGSHSYTAEEAGIGILVILIGVLLIGCMYCRNRNGYRALMDK 60
 DB 1 MPREDAFITGYGPKGSHSYTAEEAGIGILVILIGVLLIGCMYCRNRNGYRALMDK 60
 QY 61 SLHVGTCALTRRCPOEGFDHRDSKVSIOEKNCPEVVPNAPPAYEKLSAEO\$PPPYSP 118
 DB 61 SLHVGTCALTRRCPOEGFDHRDSKVSIOEKNCPEVVPNAPPAYEKLSAEO\$PPPYSP 118

RESULT 13
 AAU77810
 ID AAU77810 standard; Protein; 118 AA.

XX

AC AAU77810;
 XX
 DT 05-JUN-2002 (first entry)
 XX
 DE Human MART-1 protein mutant #5.
 XX
 KW MART-1; melanoma antigen recognised by T cells; human; mutant;
 KM anti-melanoma peptide; major histocompatibility complex; MHC;
 XX immunotherapy; cancer; vaccine; immunoregulatory; mutein.
 OS Homo sapiens.
 XX
 FT Key Location/Qualifiers
 FT Misc-difference 27 /note- "Wild type A substituted by T"
 FT Misc-difference 28 /note- "Wild type A substituted by A"
 XX
 PN WO200212272-A2.
 XX
 PD 14-FEB-2002.
 XX
 PF 03-AUG-2001; 2001WO-US24328.
 XX
 PR 04-AUG-2000; 2000US-223641P.
 PR 13-DEC-2000; 2000US-255502P.
 PR 25-JAN-2001; 2001US-264432P.
 PR 26-MAR-2001; 2001US-279005P.
 XX
 PA (GENZ) GENZYME CORP.
 XX
 PI Nicolette CA;
 XX
 DR WPI; 2002-257459/30.
 XX
 PT Novel anti-melanoma peptide compounds useful for inducing immune
 PT response in a subject, and in the preparation of medicaments for the
 PT treatment and diagnosis of cancer
 XX
 PS Claim 16; Page -; 79pp; English.
 XX
 CC This invention relates to a novel anti-melanoma peptide compound
 CC comprising a peptide of the human melanoma antigen recognised by T cells
 CC (MART-1) protein. These compounds are designed to enhance binding to
 CC major histocompatibility (MHC) compounds and enhance immunoregulatory
 CC properties and induce an immune response. The invention also comprises
 CC the nucleotide sequences encoding the peptides of the invention. The
 CC compounds of the invention are useful for inducing an immune response in
 CC a subject, by delivering the compounds in the context of a major
 CC histocompatibility (MHC) molecule which presents the compound on the
 CC surface of an antigen presenting cell, or by delivering it as a
 CC polynucleotide that encodes the peptide. The invention also comprises
 CC antibodies that recognise and bind these compounds which are useful in
 CC immunotherapy. The compounds of the invention are useful for modulating
 CC an immune response to synthetic and naturally occurring compounds in a
 CC subject. The compounds may also be used as components of anti-cancer
 CC vaccines and to expand immune effector cells that are specific for
 CC cancers characterised by expression of the human melanoma antigen
 CC recognised by T cells, MART-1. The compounds of the invention are also
 CC useful for the detection and purification of antibodies and may be used
 CC for the preparation of medicaments for the diagnosis and treatment of
 CC diseases such as cancer. The compounds of the invention have enhanced
 CC binding to MHC molecules and enhanced immunoregulatory properties
 CC relative to their natural counterparts. The present sequence represents
 CC the human melanoma antigen recognised by T cells (MART-1) mutant #5 of
 CC the invention. This mutant has amino acid alterations in the region
 CC corresponding to the MHC class I binding site, these mutations confer
 CC tighter binding to the MHC.
 CC Note: This sequence is not shown in the specification but was created by
 CC the indexer from the wild type sequence shown in AAU7793 and the
 CC information given in claim 16 of the specification.
 XX
 SO Sequence 118 AA;

Query Match 99.4%; Score 645; DB 23; Length 118;
 Best Local Similarity 99.2%; Pred. No. 66-65;
 Matches 117; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 MPRDAHFTYGYPKKGHSHSYTAEAGIGITVILGVLLIGCWYCRRRNGRYALMDK 60
 |||||
 DB 1 MPRDAHFTYGYPKKGHSHSYTAEAGIGITVILGVLLIGCWYCRRRNGRYALMDK 60
 |||||

OY 61 SLHVGTCALTRRCPOEGFDHRDSKYSLOEKNCPEVVPNAPPAVEKLSAQSPPPYSP 118
 |||||
 DB 61 SLHVGTCALTRRCPOEGFDHRDSKYSLOEKNCPEVVPNAPPAVEKLSAQSPPPYSP 118
 |||||

RESULT 14
 AAU77809.
 ID AAU77809 standard; Protein; 118 AA.
 XX
 AC AAU77809;
 XX
 DT 05-JUN-2002 (first entry)
 XX
 DE Human MART-1 protein mutant #4.
 XX
 XX MART-1; melanoma antigen recognised by T cells; human; mutant;
 KW anti-melanoma peptide; major histocompatibility complex; MHC;
 KW immunotherapy; cancer; vaccine; immunoregulatory; muteln.
 XX
 OS Homo sapiens.
 XX
 XX Key Location/Qualifiers
 FH MISC-difference 27
 FT /note= "Wild type A substituted by F"
 FT MISC-difference 28
 FT /note= "Wild type A substituted by A"
 XX
 FT WO200212272-A2.
 XX
 PN 14-FEB-2002.
 XX
 PD 03-AUG-2001; 2001WO-US24328.
 XX
 PE 04-AUG-2000; 2000US-223641P.
 PR 13-DEC-2000; 2000US-255502P.
 PR 25-JAN-2001; 2001US-264432P.
 PR 26-MAR-2001; 2001US-279005P.
 XX
 PA (GENZ) GENZYME CORP.
 XX
 PI Nicolette CA;
 XX
 DR WPI: 2002-257459/30.
 XX
 FT Novel anti-melanoma peptide compounds useful for inducing immune
 PT response in a subject, and in the preparation of medicaments for the
 PT treatment and diagnosis of cancer
 XX
 PS Claim 15; Page -: 79pp; English.
 XX
 CC This invention relates to a novel anti-melanoma peptide compound
 CC comprising a peptide of the human melanoma antigen recognised by T cells
 CC (MART-1) protein. These compounds are designed to enhance binding to
 CC major histocompatibility (MHC) compounds and enhance immunoregulatory
 CC properties and induce an immune response. The invention also comprises
 CC the nucleotide sequences encoding the peptides of the invention. The
 CC compounds of the invention are useful for inducing an immune response in
 CC a subject, by delivering the compounds in the context of a major
 CC histocompatibility (MHC) molecule which presents the compound on the
 CC surface of an antigen presenting cell, or by delivering it as a
 CC polynucleotide that encodes the peptide. The invention also comprises
 CC antibodies that recognise and bind these compounds which are useful in
 CC immunotherapy. The compounds of the invention are useful for modulating
 CC an immune response to synthetic and naturally occurring compounds in a

CC subject. The compounds may also be used as components of anti-cancer
 CC vaccines and to expand immune effector cells that are specific for
 CC cancers characterised by expression of the human melanoma antigen
 CC recognised by T cells. MART-1. The compounds of the invention are also
 CC useful for the detection and purification of antibodies and may be used
 CC for the preparation of medicaments for the diagnosis and treatment of
 CC diseases such as cancer. The compounds of the invention have enhanced
 CC binding to MHC molecules and enhanced immunoregulatory properties
 CC relative to their natural counterparts. The present sequence represents
 CC the human melanoma antigen recognised by T cells (MART-1) mutant #4 of
 CC the invention. This mutant has amino acid alterations in the region
 CC corresponding to the MHC class I binding site, these mutations confer
 CC tighter binding to the MHC.
 CC Note: This sequence is not shown in the specification but was created by
 CC the indexer from the wild type sequence shown in AAU77793 and the
 CC information given in claim 15 of the specification.
 XX
 SQ Sequence 118 AA;

Query Match 99.1%; Score 643; DB 23; Length 118;
 Best Local Similarity 99.2%; Pred. No. 1e-64;
 Matches 117; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 MPRDAHFTYGYPKKGHSHSYTAEAGIGITVILGVLLIGCWYCRRRNGRYALMDK 60
 |||||
 DB 1 MPRDAHFTYGYPKKGHSHSYTAEAGIGITVILGVLLIGCWYCRRRNGRYALMDK 60
 |||||

OY 61 SLHVGTCALTRRCPOEGFDHRDSKYSLOEKNCPEVVPNAPPAVEKLSAQSPPPYSP 118
 |||||
 DB 61 SLHVGTCALTRRCPOEGFDHRDSKYSLOEKNCPEVVPNAPPAVEKLSAQSPPPYSP 118
 |||||

RESULT 15
 AAU31980
 ID AAU31980 standard; Protein; 118 AA.
 XX
 AC AAU31980;
 XX
 DT 21-DEC-1999 (first entry)
 XX
 DE Human MART1 melanocyte differentiation antigen.
 XX
 XX MART1; melanocyte differentiation antigen; melanoma; human;
 KW antigen presentation; adoptive immunotherapy; cancer; therapy;
 KW vaccine.
 XX
 OS Homo sapiens.
 XX
 PN WO9947102-A2.
 XX
 PD 23-SEP-1999.
 XX
 PE 19-MAR-1999; 99WO-US06031.
 PR 20-MAR-1998; 98US-0078880.
 XX
 PA (GENZ) GENZYME CORP.
 XX
 PI Nicolette CA, Kaplan J;
 XX
 DR WPI: 1999-590956/50.
 DR N-PSDB; AAU20065.
 XX
 PT Preparing cells for use as cancer vaccines and in adoptive
 PT immunotherapy
 XX
 PS Disclosure; Page 51; 55pp; English.
 XX
 CC The present sequence represents human MART1, a melanocyte
 CC differentiation antigen that is specifically recognised by HLA-A2
 CC restricted tumour-infiltrating lymphocytes of melanoma patients.
 CC The invention provides methods for immunotherapy, in particular for
 CC inducing an immune response against an antigen in a patient.

CC Genetically modified antigen-presenting cells (APC) that are more
 CC potent presenters of exogenous peptide than the parental
 CC antigen-presenting cells are used. These APCs lack an effective
 CC endogenous TAP (transporter associated with antigen processing)
 CC activity and present exogenous antigen on the major
 CC histocompatibility complex class I (MHC-I) molecule. Suitable
 CC exogenous antigens include a tumour antigen, such as a minimal
 CC essential epitope of MART-1, which can complex with MHC-I. The
 CC genetically modified APCs are useful for inducing an immune response
 CC (claimed) against an antigen in a patient (adoptive immunotherapy),
 CC especially as vaccines against cancer in mammals, preferably humans.
 CC The cells are also useful for expanding populations of immune
 CC effector cells, preferably cytotoxic T lymphocyte cells.

XX
 SQ Sequence 118 AA;

Query Match 98.9%; Score 642; DB 20; Length 118;

Best Local Similarity 99.2%; Pred. No. 1.3e-64;

Matches 117; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 MPREDAHFTYGYPRKGHSHYTAEEAAGIGITVTIIGVLLIGCWYCRNRNGYRALMDK 60

Db 1 MPREDAHFTYGYPRKGHSHYTAEEAAGIGITVTIIGVLLIGCWYCRNRNGYRALMDK 60

OY 61 SLHVGTCALTRRCPOEGFDHRDSKVSIOEKNCPEVVPNAPPAYEKLAEQSPPPYSP 118

Db 61 SLHVGTCALTRRCPOEGFDHRDSKVSIOEKNCPEVVPNAPPAYEKLAEQSPPPYSP 118

Search completed: October 7, 2003, 18:48:16
 Job time : 42 secs

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OM protein - protein search, using sw model

Run on: October 7, 2003, 18:43:14 ; Search time 33 Seconds

(Without alignments) 922.734 Million cell updates/sec

Title: US-09-898-860-2

Perfect score: 649
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Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 830525 seqs, 258052604 residues

Total number of hits satisfying chosen parameters: 830525

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%

Listing first 45 summaries

Database :

1: SP. ARCHAEA: *
2: SP. BACTERIA: *
3: SP. FUNGI: *
4: SP. HUMAN: *
5: SP. INVERTEBRATE: *
6: SP. MAMMAL: *
7: SP. MHC: *
8: SP. ORGANELLE: *
9: SP. PHAGE: *
10: SP. PLANT: *
11: SP. RODENT: *
12: SP. VIRUS: *
13: SP. VERTEBRATE: *
14: SP. UNCLASSIFIED: *
15: SP. VIRUS: *
16: SP. BACTERIAP: *
17: SP. ARCHAEP: *

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	420.5	64.8	113	11	Q9DIY3
2	87	13.4	4957	4	014687
3	87	13.4	5262	4	014686
4	81	12.5	2034	2	093NX7
5	80.5	12.4	344	11	Q9R201
6	80.5	12.4	1127	3	Q9P571
7	79.5	12.2	494	15	0994J1
8	78	12.0	540	17	08TUC3
9	78	12.0	1814	10	08M5N5
10	77	11.9	488	4	08N504
11	77	11.9	491	4	08RBU7
12	77	11.9	750	10	09AUC2
13	76	11.7	303	4	09UKJ1
14	75.5	11.6	227	4	09UKJ0
15	75	11.6	107	12	09J455
16	75	11.6	107	12	086198

17	74.5	11.5	496	15	08UT44	08UT44 human immun
18	74	11.4	739	5	08IGR9	08IGR9 drosophila
19	74	11.4	883	5	09YBN1	09YBN1 drosophila
20	73.5	11.3	198	11	08R182	08R182 mus musculus
21	73	11.2	301	5	08MY26	08MY26 papilio xut
22	73	11.2	1069	11	088185	088185 mus musculus
23	72.5	11.2	393	6	08MK55	08MK55 cyclopes gl
24	72.5	11.2	405	3	08J221	08J221 glareia lozo
25	72.5	11.2	1427	5	09YPO0	09YPO0 drosophila
26	72	11.1	1729	10	08R2W5	08R2W5 oryza sativ
27	71.5	11.0	1265	2	P72316	P72316 rhodospirill
28	71	10.9	412	4	09H5E0	09H5E0 homo sapien
29	71	10.9	500	15	09IVA4	09IVA4 human immun
30	71	10.9	555	4	08MB59	08MB59 homo sapien
31	70.5	10.9	179	4	08RAS3	08RAS3 homo sapien
32	70.5	10.9	254	16	Q91024	Q91024 pseudomonas
33	70.5	10.9	348	2	052641	052641 pseudomonas
34	70.5	10.9	348	2	059715	059715 pseudomonas
35	70.5	10.9	348	2	051972	051972 pseudomonas
36	70.5	10.9	402	4	096163	096163 homo sapien
37	70.5	10.9	449	15	Q999X1	Q999X1 feline immu
38	70.5	10.9	449	15	Q999X2	Q999X2 feline immu
39	70.5	10.9	592	11	Q9JLN5	Q9JLN5 mus musculi
40	70	10.8	112	6	09GMM4	09GMM4 macaca fasc
41	70	10.8	253	6	0951B6	0951B6 ovis aries
42	70	10.8	335	5	09TXV5	09TXV5 caenorhabdi
43	70	10.8	346	10	Q9ZPT9	Q9ZPT9 arabidopsis
44	70	10.8	389	16	08PKX9	08PKX9 xanthomonas
45	70	10.8	562	16	Q9K3S5	Q9K3S5 streptomyces

ALIGNMENTS

RESULT 1	ID	Q9DIY3	PRELIMINARY:	PRT:	113 AA.
AC	Q9DIY3	01-UN-2001 (TREMBLrel. 17, Created)			
DT	01-UN-2001 (TREMBLrel. 17, Last sequence update)				
DT	01-UN-2001 (TREMBLrel. 17, Last annotation update)				
DE	A930034P04RIK				
GN	A930034P04RIK				
OS	Mus musculus (Mouse)				
OC	Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;				
OC	Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.				
OX	NCBI_TaxID=10090;				
RN	[1]				
RP	SEQUENCE FROM N. A.				
RC	STRAIN=C57BL/6J; TISSUE=Retina;				
RX	MEDLINE=21085660; PubMed=11217851;				
RA	Kawai J., Shingawa A., Shibata K., Yoshino M., Itoh M., Ishii Y.,				
RA	Arakawa T., Hara A., Fukunishi Y., Kono H., Adachi J., Fukuda S.,				
RA	Alzawa K., Izawa M., Nishi K., Kiyosawa H., Kondo S., Yamana I.,				
RA	Saito T., Okazaki Y., Gojobori T., Bono H., Kasukawa T., Saito R.,				
RA	Kadota K., Matsuda H.A., Ashburner M., Batalov S., Casavant T.,				
RA	Fleischmann W., Gaasterland T., Gissi C., King B., Kochiwa H.,				
RA	Kuehl P., Lewis S., Matsuo Y., Nikaido I., Pesole G., Quackenbush J.,				
RA	Schriml L.M., Staubli F., Suzuki R., Tomita M., Wagner R., Washio T.,				
RA	Sakel K., Okido T., Furuno M., Aono H., Baldarelli R., Barsi G.,				
RA	Blake J., Boffelli D., Bojunga N., Carninci P., de Bonaldo M.F.,				
RA	Brownstein W.J., Bult C., Fletcher C., Fujita M., Gariboldi M.,				
RA	Gustincich S., Hill D., Hofmann M., Hume D.A., Kamya M., Lee N.H.,				
RA	Lyons P., Marchionni L., Mashima J., Mazzarelli J., Mombarts P.,				
RA	Norone P., Ring B., Ringwald M., Rodriguez I., Sakamoto N.,				
RA	Sasaki H., Sato K., Schoenbach C., Seya T., Shibata Y., Storch K.-F.,				
RA	Suzuki H., Toyooka K., Wang K.H., Weltz C., Whitaker C., Wilming L.,				
RA	Wynshaw-Boris A., Yoshida K., Hasegawa Y., Kawai H., Kohsaki S.,				
RA	Hayashizaki Y.;				
RT	"Functional annotation of a full-length mouse cDNA collection.";				
DR	Nature 409:685-690(2001).				
EMBL	AK020928: BAB32259.1;				
MGD	MGI:1925086; A930034P04RIK.				

5Q SEQUENCE 113 AA; 12818 MW; AC01FC9840640E6E CRC64;

Query Match 64.8%; Score 420.5; DB 11; Length 113;
Best Local Similarity 67.8%; Pred. No. 4.3e-40;
Matches 80; Conservative 11; Mismatches 22; Indels 5; Gaps 2;

OY 1 MPREDATFVGYPKKGHSHSYTAEEAGIGLTVLLIGWYCRRRRGYBALMDK 60
DB 1 MPOEDINH--GYPRGHRSTVTAEEANGNGLVLLIGWYCRRRRGYBALMDK 58
OY 61 SLHVGTCALTRPCQEGFDRHSKVSLSQEKNCSEVVPVNPAPYEKLSAEOSSPPYSP 118
DB 59 RRHIGTQSRKSRSCSESPDHDSRLSSQEKSHQVVPVNPAPYEKLS--SPPYSP 113

RESULT 2

014687 ID .014687 PRELIMINARY; PRT; 4957 AA.
AC .014687;
DT 01-JAN-1998 (TREMBLrel. 05, Created)
DT 01-JAN-1998 (TREMBLrel. 05, Last sequence update)
DT 01-MAR-2003 (TREMBLrel. 23, Last annotation update)
DE ALR.
GN ALR.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=97388474; Pubmed=9247308;
RA Prasad R., Zhadanov A.B., Sedkov Y., Bullrich F., Druck T.,
RA Rallapalli R., Yano T., Alder H., Croce C.M., Huebner K., Mazo A.,
RA Cnaan E.;
RT *Structure and expression pattern of human ALR, a novel gene with
RT strong homology to ALI-1 involved in acute leukemia and to Drosophila
RT trithorax.*
RL Oncogene 15:549-560(1997).
CC -1 SIMILARITY: CONTAINS 1 SET DOMAIN.
EMBL: AF010404; AAC51735.1; -
DR InterPro: IPR003889; Fitch.C.
DR InterPro: IPR003888; Fitch.N.
DR InterPro: IPR000910; HMG_12_box.
DR InterPro: IPR003616; PostSET.
DR InterPro: IPR006118; Recombinase.
DR InterPro: IPR001214; SET.
DR InterPro: IPR001965; Znf_PHD.
DR Pfam: PF00628; PHD; 3.
DR SMART: SM00542; FYRC; 1.
DR SMART: SM00541; FYRN; 1.
DR SMART: SM00398; HMG; 1.
DR SMART: SM00249; PHD; 4.
DR SMART: SM00508; PostSET; 1.
DR SMART: SM00317; SET; 1.
DR PROSITE: PS00398; RECOMBINASES_2; 1.
DR PROSITE: PS50280; SET; 1.
DR PROSITE: PS50016; ZF_PHD_2; 1.
SQ SEQUENCE 4957 AA; 531840 MW; 1026562E1419CE8D CRC64;

Query Match 13.4%; Score 87; DB 4; Length 4957;
Best Local Similarity 31.2%; Pred. No. 3.2;
Matches 25; Conservative 7; Mismatches 30; Indels 18; Gaps 3;

OY 41 LLIGCWGRRRNGRYALMDKSLHVGTCALTR---RCPQEGFDRHSKVSLSQEKNCSEPV 96
DB 939 LIIICRHCER-----WMHAGCESLFTEDVDVNAHDEGFD---CVSCQPYVVKRY 984
OY 97 VPNPAPYEKLSAEOSSPPY 116
DB 985 APVAPPPELVPMKVEPEPOY 1004

RESULT 3

014686 ID .014686 PRELIMINARY; PRT; 5262 AA.
AC .014686;
DT 01-JAN-1998 (TREMBLrel. 05, Created)
DT 01-JAN-1998 (TREMBLrel. 05, Last sequence update)
DT 01-MAR-2003 (TREMBLrel. 23, Last annotation update)
DE ALR.
GN ALR.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=97388474; Pubmed=9247308;
RA Prasad R., Zhadanov A.B., Sedkov Y., Bullrich F., Druck T.,
RA Rallapalli R., Yano T., Alder H., Croce C.M., Huebner K., Mazo A.,
RA Cnaan E.;
RT *Structure and expression pattern of human ALR, a novel gene with
RT strong homology to ALI-1 involved in acute leukemia and to Drosophila
RT trithorax.*
RL Oncogene 15:549-560(1997).
CC -1 SIMILARITY: CONTAINS 1 SET DOMAIN.
EMBL: AF010403; AAC51734.1; -
DR Gene; HGNC:7133; M12.
DR InterPro: IPR003889; Fitch.C.
DR InterPro: IPR003888; Fitch.N.
DR InterPro: IPR000910; HMG_12_box.
DR InterPro: IPR003616; PostSET.
DR InterPro: IPR006118; Recombinase.
DR InterPro: IPR001214; SET.
DR InterPro: IPR001965; Znf_PHD.
DR InterPro: IPR001841; Znf_Ring.
DR Pfam: PF00628; PHD; 5.
DR SMART: SM00542; FYRC; 1.
DR SMART: SM00541; FYRN; 1.
DR SMART: SM00398; HMG; 1.
DR SMART: SM00249; PHD; 7.
DR SMART: SM00508; PostSET; 1.
DR SMART: SM00184; RING; 3.
DR PROSITE: PS00398; RECOMBINASES_2; 1.
DR PROSITE: PS50280; SET; 1.
DR PROSITE: PS50016; ZF_PHD_2; 1.
DR PROSITE: PS50089; ZF_RING_2; 1.
SQ SEQUENCE 5262 AA; 564171 MW; 26B7C74CAD417E44 CRC64;

Query Match 13.4%; Score 87; DB 4; Length 5262;
Best Local Similarity 31.2%; Pred. No. 3.5;
Matches 25; Conservative 7; Mismatches 30; Indels 18; Gaps 3;

OY 41 LLIGCWGRRRNGRYALMDKSLHVGTCALTR---RCPQEGFDRHSKVSLSQEKNCSEPV 96
DB 1244 LIIICRHCER-----WMHAGCESLFTEDVDVNAHDEGFD---CVSCQPYVVKRY 1289
OY 97 VPNPAPYEKLSAEOSSPPY 116
DB 1290 APVAPPPELVPMKVEPEPOY 1309

RESULT 4

093NK7 ID .093NK7 PRELIMINARY; PRT; 2034 AA.
AC .093NK7;
DT 01-DEC-2001 (TREMBLrel. 19, Created)
DT 01-DEC-2001 (TREMBLrel. 19, Last sequence update)
DT 01-MAR-2003 (TREMBLrel. 23, Last annotation update)
DE AMPK.
GN AMPK.
OS Streptomyces nodosus.
OC Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;


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OC Streptomycinae; Streptomycetaceae; Streptomycetes.
OX NCBI_TaxID=40318;
RN [1]
RP SEQUENCE FROM N.A.
RA Caffrey P., Lynch S.V., Flood E.M., Finnan S.M., O'Leary M.;
RT "The amphotericin biosynthetic gene cluster from Streptomycetes
   nodosus."
RL Submitted (MAR-2001) to the EMBL/GenBank/DBJ databases.
CC -1- SIMILARITY: BELONGS TO THE SHORT-CHAIN DEHYDROGENASIS/REDUCTASES
   (SDR) FAMILY.
DR EMBL: AF357202; AAK73503.1;
DR InterPro: IPR001227; Ac_transferase.
DR InterPro: IPR002198; ADH_short.
DR InterPro: IPR000794; Ketoacyl-synt.
DR InterPro: IPR006163; Pp_bind.
DR InterPro: IPR000379; Ser_estrs_site.
DR InterPro: IPR001031; Thioesterase.
DR Pfam: PF00698; Acyl_transf. 1.
DR Pfam: PF00106; adh_short. 1.
DR Pfam: PF02801; ketoacyl-synt.C; 1.
DR Pfam: PF00550; pp-binding. 1.
DR Pfam: PF00975; Thioesterase; 1.
DR PROSITE: PS50075; ACP_DOMAIN. 1.
DR PROSITE: PS00606; B_KETOACYL_SYNT_HASE; 1.
DR Oxidoreductase; Phosphopantetheine; Transferase.
KM SEQUENCE 2034 AA; 212063 MW; 83B962B7DC3D5747 CRC64;

Query Match 12.5%; Score 81; DB 2; Length 2034;
Best Local Similarity 26.8%; Pred. NO. 5.6;
Matches 33; Conservative 14; Mismatches 44; Indels 32; Gaps 6;

OY 11 GYPRKHGSHYTTAEAGAGIGILTVILGVLILGICWCRNGRNGRADMKSLSHY----- 64
DB 370 GYVKSNIH---TQAAGAGAVIKMVLAL-----RHG---LLPRLHLDADSTH 412
OY 65 -----GTGCALTRRCP-QEGFDRHDSKVS---LQEKNCPEVVPNAPAYEKLSAQSPPP 115
DB 413 VMDADGHVSLTETATPWECEGQTRRACVSSFGISGTNAHYTLERAPAEEDTDEQREPP 472
OY 116 YSP 118
DB 473 VVP 475

RESULT 5
OY 09R201 PRELIMINARY; PRT; 344 AA.
AC 09R201;
DT 01-MAY-2000 (TREMBlrel. 13, Created)
DT 01-MAY-2000 (TREMBlrel. 13, Last sequence update)
DT 01-MAR-2003 (TREMBlrel. 23, Last annotation update)
DE T-cell surface antigen CD2.
GN CD2.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=BA1B/CJ; TISSUE=Spleen;
RA Ma R.2., Teuscher C.;
RL Submitted (MAY-1998) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=BA1B/CYJ; TISSUE=Spleen;
RA Ma R.2., Cory T.;
RT "CD2 is a candidate gene for Tmewd3 in mice."
RL Submitted (SEP-2000) to the EMBL/GenBank/DBJ databases.
DR EMBL: AF065309; AAD25889.1;
DR EMBL: AF306543; AAG27722.1;
DR HSP: P08921; IAG4.
DR MGD; MGI:88320; Cd2.

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DR InterPro: IPR000719; Prot_Kinase.
DR PROSITE: PS00107; PROTEIN_KINASE_ATP; 1.
SQ SEQUENCE 344 AA; 38325 MW; 3808BBF9FE2FE3B CRC64;

Query Match 12.4%; Score 80.5; DB 11; Length 344;
Best Local Similarity 27.4%; Pred. NO. 0.83;
Matches 31; Conservative 15; Mismatches 46; Indels 21; Gaps 5;

OY 3 REDAHFICYPRKKGSHYTTAEAGAGIGILTVILGVLILGICWCRNGRNGRADMKSLS 62
DB 188 KESKTEVNCPEKGLSF-YTVGVAG-GLLVL-LVALFIC-ICKRRRRRRRDEEL 243
OY 63 HVGTCALTRRCPQEGFDRHDSKVSLOEKNCPEVVPNAPAYEKLSAQSPPP 115
DB 244 EI-----KASRTSTVERGPFPHSTPAAAGNVALQHPPP 279

RESULT 6
OY 09P571 PRELIMINARY; PRT; 1127 AA.
AC 09P571;
DT 01-OCT-2000 (TREMBlrel. 15, Created)
DT 01-DEC-2001 (TREMBlrel. 19, Last sequence update)
DT 01-OCT-2002 (TREMBlrel. 22, Last annotation update)
DE Related to actin-interacting protein AIP3.
GN B9J10.100.
OS Neurospora crassa.
OC Eukaryota; Fungi; Ascomycota; Pezizomycotina; Sordariomycetes;
OC Sordariales; Sordariaceae; Neurospora.
OX NCBI_TaxID=5141;
RN [1]
RP SEQUENCE FROM N.A.
RA Schulte U., Algn V., Hohelsel J., Brandt P., Fartmann B., Holland R.,
RA Nykatura G., Meves H.W., Mannhaupt G.;
RL Submitted (MAY-2000) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RA German Neurospora genome project;
RL Submitted (NOV-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL: AL356324; CAB92017.2;
DR InterPro: IPR005613; AIP3.
DR Pfam: PF03915; AIP3; 1.
SQ SEQUENCE 1127 AA; 122731 MW; B6727C2CFB031DDC CRC64;

Query Match 12.4%; Score 80.5; DB 3; Length 1127;
Best Local Similarity 25.6%; Pred. NO. 3.2;
Matches 34; Conservative 16; Mismatches 52; Indels 31; Gaps 7;

OY 11 GY-PRKHGSHYTTAEAGAGIGILTVILGVLILGICWCR-----RRNGR--ALMKS 61
DB 223 GYGRKRGKPS-----SSGVPTRTSTSTYASVNSGLTNMNEGLESNGTRPDSFRDS 276
OY 62 LHVQ-TGCALTRRCPQEGFDRHDSKVSLOE-----RNC-----EPVVPNAPAYE 105
DB 277 RNSGPTASTPRRLPSQDQSGSVTSQSSLSNWTMNIPIVMADYPGERTIPGPAPP 336
OY 106 KLSAQSPPTSP 118
DB 337 EINVDNFPPTPP 349

RESULT 7
OY 0994J1 PRELIMINARY; PRT; 494 AA.
AC 0994J1;
DT 01-JUN-2001 (TREMBlrel. 17, Created)
DT 01-JUN-2001 (TREMBlrel. 17, Last sequence update)
DT 01-MAR-2003 (TREMBlrel. 23, Last annotation update)
DE Gag protein (Gag polypeptide).
GN GAG.
OS Human immunodeficiency virus 1.
OC Viruses; Retroid viruses; Retroviridae; Lentivirus.
OX NCBI_TaxID=11676;

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RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-98IN022;
RX MBLIN-21094715; PubMed-11177395;
RA Rodenburg C.M., Li Y., Trisk S.A., Chen Y., Decker J., Robertson D.L.,
RA Kalish M.L., Shaw G.M., Allen S., Hahn B.H., Gao F.;
RT "Near full-length clones and reference sequences for subtype C
RT isolates for HIV type 1 from three different continents.";
RL AIDS Res. Hum. Retroviruses 17:161-168(2001).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN-98IN022;
RA Rodenburg C.M., Li Y., Trisk S.A., Chen Y., Decker J., Robertson D.L.,
RA Allen S., Shaw G.M., Hahn B.H., Gao F.;
RT Submitted (JUL-2000) to the EMBL/GenBank/DDBJ databases.
RL EMBL: AF286232; AAK31035.1;
DR HSBP; P03888; IAA.
DR InterPro: IPR000721; Gag_P24.
DR InterPro: IPR000071; Retrovir_P17.
DR InterPro: IPR001878; Znf_CCHC.
DR Pfam: PF00540; Gag_P17; 1.
DR Pfam: PF00607; Gag_P24; 1.
DR Pfam: PF00098; Zf_CCHC; 2.
DR PRINTS; PR00939; C2HCZNFINGER.
DR SMART; SM00343; Znf_C2HC; 2.
DR PROSITE; PS50158; Zf_CCHC; 2.
KW AIDS, Core protein, Polyprotein.
SQ SEQUENCE 494 AA; 55092 MW; A2C66A4EA3DB5E35 CRC64;

Query Match 12.2%; Score 79.5; DB 15; Length 494;
Best Local Similarity 21.7%; Pred. No. 1.6;
Matches 28; Conservative 13; Mismatches 49; Indels 39; Gaps 4;

OY 16 GHGSHYTAEEAGIGLITVIL-----GVLLIGCYCRRRNGYRALMDKSLHVGTOCA 69
DB 352 GPGHKARVLAENASQANSTIMONGNFKGPKRIYKFCNC-----GREGHLAKNCR 401
OY 70 LTR-----RCPOGEF-----DHRDSTKSLQKCEPVYPNAPAEK 106
DB 402 APRKKGCGKCGKCHOKKDCITERQANFLGRIMPSHKGRPGNFIQSRPEPTVTPAPASFS 461
OY 107 LSAEQSPPP 115
DB 462 GFGETTPAP 470

RESULT 8
O8TUC3 PRELIMINARY; PRT; 540 AA.
ID O8TUC3
AC O8TUC3;
DT 01-JUN-2002 (TREMblrel. 21, Created)
DT 01-JUN-2002 (TREMblrel. 21, Last sequence update)
DT 01-OCT-2002 (TREMblrel. 22, Last annotation update)
DE Methylanine methyltransferase corrinoid activation protein.
GN RWA OR MA0150.
OS Methanosarcina acetivorans.
OC Archaea; Euryarchaeota; Methanococci; Methanosarcinales;
OC Methanosarcinaceae; Methanosarcina.
OX NCBI_TaxID=2214;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-C2A / ATCC 35395 / DSM 2834;
RX MEDLINE-21929760; PubMed-11932238;
RA Galagan J.E., Nisbaum C., Roy A., Enditzel M.G., Macdonald P.,
RA Fitzhugh W., Calvo S., Engels R., Smirnov S., Alnoor D., Brown A.,
RA Linton L., McEwen P., McKernan K., Talamas J., Tittell A., Ye W.,
RA Zimmer A., Barber R.D., Cann I., Graham D.E., Grahame D.A., Guss A.M.,
RA Hedderich R., Ingram-Smith C., Kuetner H.C., Krywcki J.A.,
RA Leight J.A., Li W., Liu J., Mukhopadhyay B., Reeve J.N., Smith K.,
RA Springer T.A., Umeyam L.A., White O., White R.H., de Macario E.C.,
RA Ferry J.G., Jarrell K.F., Jing H., Macario A.J.L., Paulsen I.,

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RA Pritchett M., Sowers K.R., Swanson R.V., Zinder S.H., Lander E.,
RA Metcalf W.W., Birren B.;
RT "The genome of Methanosarcina acetivorans reveals extensive metabolic
RT and physiological diversity.";
RL Genome Res. 12:532-542(2002).
DR EMBL; AE010672; AAM03603.1;
DR InterPro: IPR001450; 4Fe4S_ferredoxin.
DR Pfam: PF00037; fer4; 2.
DR PROSITE; PS00198; 4Fe4S_FERREDOXIN; 1.
KW Transferase; Methyltransferase; Complete proteome.
SQ SEQUENCE 540 AA; 58864 MW; 7B3D323035D5227B CRC64;

Query Match 12.0%; Score 78; DB 17; Length 540;
Best Local Similarity 26.3%; Pred. No. 2.7;
Matches 30; Conservative 17; Mismatches 31; Indels 36; Gaps 8;

OY 9 IYGPBKRGHSGYTAA-----BAAGIGLTVI--LGVLILIG--CWYCR----- 50
DB 432 IYGLPKLPRTLEHADIEKRVSDIQVGGALAIKEIILILEVSDKILYQCKVECP 491
OY 51 -----RNGR-ALMDKSLHVGTOCALTR-----CPQEGDHRDSKVSLOK 91
DB 492 EGALIEVTNGNRIRAKYDSQKCIQSGC---RRCVSICPNALD--ITKIKER 540

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RESULT 9
O8W5N5 PRELIMINARY; PRT; 1814 AA.
ID O8W5N5
AC O8W5N5;
DT 01-MAR-2002 (TREMblrel. 20, Created)
DT 01-MAR-2002 (TREMblrel. 20, Last sequence update)
DT 01-MAR-2003 (TREMblrel. 23, Last annotation update)
DE Putative disease resistant protein.
GN OSUNBA001K12.15.
OS Oryza sativa (Rice).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
OC Ehrhartoideae; Oryzaceae; Oryza.
OX NCBI_TaxID=4530;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-cv. Nipponbare;
RA Buell C.R., Yuan Q., Ouyang S., Liu J., Moffat K.S., Hill J.N.,
RA Ganabergner K., Brenner M., Burgess S., Hance M., Shwartsbeyn M.,
RA Tsirlin T., Riggs F., Hsiao J., Zismann V., Blunt S., Pal G.,
RA Vanden S.E., Utterback T.R., Feldblyum T.V., Raib E., Quackenbush J.,
RA Salzberg S.L., White O., Fraser C.M.;
RT "Oryza sativa chromosome 10 BAC OSUNBA0001K12 genomic sequence.";
RL Submitted (NOV-2001) to the EMBL/GenBank/DDBJ databases.
DR EMBL; AC078948; AAL31031.1;
DR Gramene; O8W5N5;
DR InterPro: IPR000767; Disease_resist.
DR InterPro: IPR001611; LRR.
DR InterPro: IPR002182; NB-ARC.
DR Pfam; PF00560; LRR; 2.
DR Pfam; PF00931; NB-ARC; 1.
DR PRINTS; PR00364; DISEASERISST.
SQ SEQUENCE 1814 AA; 19893 MW; C18A0AED6C4AD27 CRC64;

Query Match 12.0%; Score 78; DB 10; Length 1814;
Best Local Similarity 27.5%; Pred. No. 11;
Matches 28; Conservative 11; Mismatches 33; Indels 30; Gaps 5;

OY 30 IGLTVILGVLLIGCYCRRRNGYRALMDKSLHVGTOCALTRRCPQ----- 76
DB 1599 VAILLITISVAOULFTFM-----YRKLMNSKYGKETSNSDCPQFSFSLTLTFL 1652
OY 77 EGFDHRDSKVSLOKCEPVYPNAPAEKLSABQSPPPYP 118
DB 1653 ESF-----SVDEEVKIPSP--PP--EKEEFAISPQPP 1683

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RESULT 10

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OBN504
ID OBN504 PRELIMINARY: PRT; 488 AA.
AC OBN504:
DT 01-OCT-2002 (TREMBLrel. 22, Created)
DT 01-OCT-2002 (TREMBLrel. 22, Last sequence update)
DT 01-MAR-2003 (TREMBLrel. 23, Last annotation update)
DE Hypothetical protein (Fragment).
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Melanoma;
RA Strausberg R.;
RL Submitted (JUN-2002) to the EMBL/GenBank/DBJ databases.
DR EMBL; BC033132; AAH3132.1; -.
DR InterPro: IPR006586; ADAM_cysteine.
DR InterPro: IPR001762; Disintegrin.
DR InterPro: IPR006209; EGF like.
DR Pfam; PF00200; disintegrin.1.
DR PRINTS; PR00289; DISINTEGRIN.
DR ProDom; PD000664; Disintegrin; 1.
DR SMART; SM00608; ACR; 1.
DR PROSITE; PS50214; DISINTEGRIN_2; 1.
DR PROSITE; PS01186; EGF_2; 1.
FW Hypothetical protein.
KW NON_TER
SQ SEQUENCE 488 AA; 52722 MW; F9D288B23529BC05 CRC64;

Query Match 11.9%; Score 77; DB 4; Length 488;
Best Local Similarity 25.5%; Pred. No. 3.1;
Matches 35; Conservative 16; Mismatches 54; Indels 32; Gaps 7;

QY 4 EDNAFTGYGP-----KKGHGSHYTTA---EEAG---IGILTVILGVLILGICWCYCRRN 52
DB 238 QNCHCLPGWAPPCNTPGHGSIDSGMPPEVGPVAGVLAIVLAVLMLMYCCRON 297
QY 53 GYRALMDKSLHVTGQALTR-----CP-----QEGFDHRSKYSIQEKNCPEVVPNAP 101
DB 298 NKILQQLKPS-----ALPSKLRQGFSCFPRVSONSGTHANPFLQTPGCKRKYINTP 350
QY 102 PAYEKLSAEQSPPPSP 118
DB 351 ---ELLRKPSQPPPPPP 364

RESULT 11
O8TBUT7 PRELIMINARY: PRT; 491 AA.
AC O8TBUT7:
DT 01-JUN-2002 (TREMBLrel. 21, Created)
DT 01-JUN-2002 (TREMBLrel. 21, Last sequence update)
DT 01-MAR-2003 (TREMBLrel. 23, Last annotation update)
DE Hypothetical protein (Fragment).
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=uterus;
RA Strausberg R.;
RL Submitted (FEB-2002) to the EMBL/GenBank/DBJ databases.
DR EMBL; BC024214; AAH24214.1; -.
DR InterPro: IPR006586; ADAM_cysteine.
DR InterPro: IPR001762; Disintegrin.
DR InterPro: IPR006209; EGF like.
DR Pfam; PF00200; disintegrin.1.
DR ProDom; PD000664; Disintegrin; 1.
DR SMART; SM00608; ACR; 1.
DR SMART; SM00050; DISIN; 1.

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DR PROSITE; PS50214; DISINTEGRIN_2; 1.
DR PROSITE; PS01186; EGF_2; 1.
KW Hypothetical protein.1
FW NON_TER
SQ SEQUENCE 491 AA; 53045 MW; BACD338C38496A08 CRC64;

Query Match 11.9%; Score 77; DB 4; Length 491;
Best Local Similarity 25.5%; Pred. No. 3.1;
Matches 35; Conservative 16; Mismatches 54; Indels 32; Gaps 7;

QY 4 EDNAFTGYGP-----KKGHGSHYTTA---EEAG---IGILTVILGVLILGICWCYCRRN 52
DB 241 QNCHCLPGWAPPCNTPGHGSIDSGMPPEVGPVAGVLAIVLAVLMLMYCCRON 300
QY 53 GYRALMDKSLHVTGQALTR-----CP-----QEGFDHRSKYSIQEKNCPEVVPNAP 101
DB 301 NKILQQLKPS-----ALPSKLRQGFSCFPRVSONSGTHANPFLQTPGCKRKYINTP 353
QY 102 PAYEKLSAEQSPPPSP 118
DB 354 ---ELLRKPSQPPPPPP 367

RESULT 12
O9AUC2 PRELIMINARY: PRT; 750 AA.
AC O9AUC2:
DT 01-JUN-2001 (TREMBLrel. 17, Created)
DT 01-JUN-2001 (TREMBLrel. 17, Last sequence update)
DT 01-MAR-2003 (TREMBLrel. 23, Last annotation update)
DE Receptor-like protein kinase 1.
FW PKRL
GN Zea mays (Maize).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
OC PACAD clade; Panicoideae; Andropogoneae; Zea.
OX NCBI_TaxID=4577;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=cv. B73;
RA Kim H.-U., Colter R., McCormick S.;
RT Arabidopsis: the tomato kinases identified in tomato, maize and
RT expression patterns during pollen tube growth."
RL Submitted (Apr-2001) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=cv. B73;
RA Dodds P., Kulikauskas R.;
RL Submitted (MAR-2000) to the EMBL/GenBank/DBJ databases.
DR EMBL; AF243041; AAK28346.1; -.
DR InterPro: IPR001611; LRR.
DR InterPro: IPR007090; LRR_plant.
DR InterPro: IPR00719; Pro_kinase.
DR InterPro: IPR002965; P_rich_extensn.
DR Pfam; PF000560; LRR; 4.
DR Pfam; PF00069; pkinase; 1.
DR PRINTS; PR01217; PRICHEXTENS.
DR ProDom; PD000001; ProL_kinase; 1.
DR PROSITE; PS50011; PROTEIN_KINASE_DOM; 1.
DR ATP-binding; kinase; transferase.
KW SEQUENCE 750 AA; 79386 MW; 34B12797F000351B CRC64;

Query Match 11.9%; Score 77; DB 10; Length 750;
Best Local Similarity 24.3%; Pred. No. 5.1;
Matches 25; Conservative 14; Mismatches 40; Indels 24; Gaps 3;

QY 14 KKGHGSHYTTAEAGIGILTVILGVLILGICW---CRNRGTRALMDKSLHVTGQALT 71
DB 305 KQEGHKNPVGSGTSTFGLVLAIFIGTLGAGVAFVALRRRGY-----KTKNGKPTAS-- 357
QY 72 RRCPEGFDHRSKYSIQEKNCPEVVPNAPPAYEKLSAEQSP 114

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Db      358 -----SARPSDPPEVEPHHPAAKAEASAAQAP 385
          | : : | : | : | : |

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RESULT 13

ID	SEQUENCE	303 AA:	33877 MW:	0410ADFC7E80928B CRC64:
AC	09UKJ1	PRELIMINARY;	PRF;	303 AA.
DT	01-MAY-2000 (TREMBLrel. 13, Created)			
DT	01-MAY-2000 (TREMBLrel. 13, Last sequence update)			
DT	01-MAR-2003 (TREMBLrel. 23, Last annotation update)			
DE	Inhibitory receptor PIRAlpha.			
OS	Homo sapiens (Human).			
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;			
OC	Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.			
OX	NCBI_TaxID=9606;			
RN	[1]			
RP	SEQUENCE FROM N.A.			
RX	MEDLINE=20127940; PubMed=10660620;			
RA	Mousseau D.D., Banville D., L'Abbe D., Bouchard P., Shen S.H.;			
RT	"PIRAlpha, a novel immunoreceptor tyrosine-based inhibitory motif-			
RT	bearing protein, recruits SHP-1 upon tyrosine phosphorylation and is			
RT	paired with the truncated counterpart PIRbeta.";			
RL	J. Biol. Chem. 275:4467-4474(2000).			
DR	EMBL: AF161080; AAD52964.1; "			
DR	InterPro: IPR003599; I9.			
DR	InterPro: IPR007110; I9-like.			
DR	SMART: SM00409; IG; 1.			
DR	PROSITE: PSS0835; IG_LIKE; 1.			
KW	Receptor.			

RESULT 14

ID	PRELIMINARY:	PRT:	227 AA.
AC	09UKJ0:		
DT	01-MAY-2000 (TREMBLrel. 13. Created)		
DT	01-MAY-2000 (TREMBLrel. 13. Last sequence update)		
DT	01-MAR-2003 (TREMBLrel. 23. Last annotation update)		
DE	Activating receptor PILRBeta.		
OS	Homo sapiens (Human).		
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;		
OC	Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.		
CC	NCBI_TaxId=9606;		
RN	[1]		
RP	SEQUENCE FROM N.A.		
RA	MEDLINE:20127940; PubMed=10660620;		
RT	Mousseau D.D., Baivillie D., V'Abbe D., Bouchard P., Shen S.H.;		
RT	PilRBeta, a novel immunoreceptor tyrosine-based inhibitory motif-		
RT	bearing protein, recruits SHP-1 upon tyrosine phosphorylation and is		
RT	paired with the truncated counterpart PILRBeta.*;		
RL	J. Biol. Chem. 275:4467-4474(2000).		
DR	EMBL: AF161081; AAD52965.1; --		
DR	InterPro: IPR003599; I9.		
DR	InterPro: IPR007110; I9-like.		
DR	SMART: SM00409; IG; 1.		
DR	PROSITE: PS50835; IG_LIKE; 1.		
KW	Receptor.		
SO	SEQUENCE	227 AA; 25542 MW; 7FF960C60AB7EFD9 CRC64;	

Query Match	11.6%;	Score 75.5;	DB 4;	Length 227;
Best Local Similarity	43.8%;	Pred. No. 1.9;		
Matches 21; Conservative	6;	Mismatches 14;	Indels 7;	Gaps 3;

RESULT 15

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ID      09J455;
AC      09J455;
AD      PRELIMINARY;
DT      01-OCT-2000 (TREMBLrel. 15, Created)
DT      01-OCT-2000 (TREMBLrel. 15, Last sequence update)
DT      01-JUN-2001 (TREMBLrel. 17, Last annotation update)
DE      Putative fusion protein (Nonstructural protein).
OS      Human rotavirus B.
OC      Viruses; dsRNA viruses; Reoviridae; Rotavirus.
OX      NCBI_TaxID:10942;
RN      [1]
RP      SEQUENCE FROM N.A.
RC      STRAIN=CA1;
RA      Sen A., Das S., Krishnan T., Kobayashi N., Nalk T.N.;
RT      "partial sequence of cDNA of the gene segment 6 encoding fusion
RL      protein and a nonstructural protein of the human group B rotavirus
RT      strain, CAL-1."
RL      Submitted (NOV-1999) to the EMBL/Genbank/DBSJ databases.
RN      [2]
RP      SEQUENCE FROM N.A.
RC      STRAIN=CA1;
RA      Kobayashi N., Nalk T.;
RT      "Sequence of the sixth RNA segment of group B human rotavirus CAL."
RT      Submitted (FEB-2000) to the EMBL/Genbank/DBSJ databases.
DR      EMBL; AF203975; AAF69263.1; -
DR      EMBL; AF230975; AAG09741.1; -
SO      SEQUENCE 107 AA; 11697 MW; 3AD1556D92B387FE CRC64;

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Search completed: October 7, 2003, 18:49:15
Job time : 35 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: October 7, 2003, 18:42:34 ; Search time 11 Seconds

(without alignments)
504.468 Million cell updates/sec

Title: US-09-898-860-2

Sequence: 1 MPREDAHFTGYGPKKGHS.....NAPPAYEKLSAEQSPPPYSP 118

Scoring table:

BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 127863 seqs, 47026705 residues

Total number of hits satisfying chosen parameters: 127863

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database: SwissProt_41:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	64.9	100.0	118	MAR1_HUMAN	Q16655 homo sapien
2	80.5	12.4	344	CD2_MOUSE	P08920 mus musculus
3	77	11.9	710	NASC_BACSU	P42434 bacillus su
4	77	11.9	956	AD15_MOUSE	Q9h013 homo sapien
5	72.5	11.2	815	AD15_MOUSE	O88839 mus musculus
6	72.5	11.2	1429	EXPA_DROME	Q07436 drosophila
7	71	10.9	1069	PCHT7_HUMAN	O60245 homo sapien
8	70.5	10.9	202	TMG2_HUMAN	O14669 homo sapien
9	69.5	10.7	218	TMG1_HUMAN	O14668 homo sapien
10	69.5	10.7	401	LSG1_HAEN	P13399 haemophilus
11	68	10.5	165	CYTL_MOUSE	O6v1b7 mus musculus
12	67.5	10.4	188	COME_MERYA	P58416 methanococc
13	67.5	10.4	778	MGD1_HUMAN	Q9y5v3 homo sapien
14	67.5	10.4	4590	FATH_HUMAN	Q14517 homo sapien
15	66.5	10.2	775	MGD1_MOUSE	Q9qyh6 mus musculus
16	66.5	10.2	775	MGD1_MOUSE	O9es73 rattus norv
17	66	10.2	140	YD23_YEAST	O67549 saccharomyc
18	66	10.2	969	SACB_STRLS	Q55242 streptococc
19	65.5	10.1	258	BOX3_MOTVI	Q95681 streptococc
20	65.5	10.1	659	SYT_THETH	P53771 notophthalm
21	65.5	10.1	1436	WC11_BOVIN	P56881 thermus the
22	65.5	10.0	309	YHCC_ECOLI	P30205 bos taurus
23	65	10.0	356	RS41_ARATH	P85450 escherichia
24	64.5	9.9	154	CYTL_HUMAN	P92966 arabidopsis
25	64.5	9.9	359	FIX2_RHILE	Q96786 homo sapien
26	64	9.9	144	TIM2_RABIT	P07748 rhizobium 1
27	64	9.9	344	CD2_RAT	O9trz7 oryzocolagus
28	64	9.9	352	CD5L_MOUSE	P08921 rattus norv
29	64	9.9	2180	POLG_ECH2H	O6578 mus musculus
30	63.5	9.8	118	VPA_STYAI	O6578 e genome po
31	63.5	9.8	227	NK1L_MOUSE	Q02842 simian immu
32	63.5	9.8	628	GIDA_WIGBR	P27811 mus musculus
33	63.5	9.8	1338	ACIN_MOUSE	O8d3k0 wigglewort
					Q9j1x8 mus musculus

ALIGNMENTS

34	63	9.7	117	1	HYPA_HELPY	O25539 helicobacte
35	63	9.7	336	1	EPB2_MOUSE	P52800 mus musculus
36	63	9.7	387	1	GIR_HUMAN	O9n183 homo sapien
37	63	9.7	629	1	GIDA_HAEN	P44763 haemophilus
38	63	9.7	1004	1	PHC1_HUMAN	P78364 homo sapien
39	63	9.7	1012	1	PHC1_MOUSE	O64028 mus musculus
40	62.5	9.6	757	1	KNC3_HUMAN	O14803 homo sapien
41	62.5	9.6	786	1	CRJ2_HUMAN	O9nq75 homo sapien
42	62.5	9.6	821	1	ENV_STYGB	P22380 simian immu
43	62.5	9.6	889	1	KNC3_RAT	O01956 rattus norv
44	62.5	9.6	913	1	DDRL_HUMAN	O08345 h epithelia
45	62	9.6	286	1	TESB_HAEN	P44498 haemophilus

RESULT 1

ID	MAR1_HUMAN	STANDARD:	PRT:	118 AA.
AC	Q16655			
DT	01-NOV-1997 (Rel. 35, Created)			
DT	01-NOV-1997 (Rel. 35, Last sequence update)			
DT	15-SEP-2003 (Rel. 42, Last annotation update)			
DE	Melanoma antigen recognized by T-cells 1 (MART-1) (Melan-A protein)			
DE	(Antigen SK29-AA) (Antigen LB39-AA).			
CN	MLANA OR MART1.			
OS	Homo sapiens (Human).			
OC	Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;			
OC	Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.			
OX	NCBI_Taxid=9606;			
RN	[1]			
RP	SEQUENCE FROM N.A.			
RC	TISSUE=Skin;			
RX	MEDLINE=942275389; PubMed=8170938;			
RX	MEDLINE=94224770; PubMed=8170938;			
RA	Kawakami Y., Ellyhu S., Delgado C.H., Robbins P.F., Rivoltini L.,			
RA	Topalian S.L., Miki T., Rosenberg S.A.;			
RT	*Cloning of the gene coding for a shared human melanoma antigen			
RT	recognized by autologous T cells infiltrating into tumor.*;			
RL	Proc. Natl. Acad. Sci. U.S.A. 91:3515-3519(1994).			
RN	[2]			
RP	SEQUENCE FROM N.A.			
RC	TISSUE=Skin;			
RX	MEDLINE=22388257; PubMed=12477932;			
RA	Straussberg R.L., Felingold E.A., Grouse L.H., Derge J.G.,			
RA	Klausner R.D., Collins F.S., Wagner L., Shemen C.M., Schuler G.D.,			
RA	Altschul S.F., Zeeberg B., Buetow K.H., Scheffer C.F., Bhat N.K.,			
RA	Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hong L.,			
RA	Ditchevko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,			
RA	Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,			
RA	Brownstein M.J., Usdin T.B., Toshiyuki S., Carinci P., Prange C.,			
RA	Rata S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullighy S.J.,			
RA	Bozak S.A., McEwan P.J., McKernan K.O., Malek J.A., Gunaratne P.H.,			
RA	Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,			
RA	Vallatton D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,			
RA	Falley J., Helton E., Kettman M., Madan A., Rodriguez S., Sanchez A.,			
RA	Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,			
RA	Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,			
RA	Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,			
RA	Butterfield Y.S.N., Krzyzinski M.I., Skalska U., Smalhus D.E.,			
RA	Schuerch A., Schein J.E., Jones S.J.M., Marra M.A.;			
RT	*Generation and initial analysis of more than 15,000 full-length			
RT	human and mouse cDNA sequences.*;			
RL	Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).			

CC -1- TISSUE SPECIFICITY: EXPRESSION IS RESTRICTED TO MELANOMA AND
 CC MELANOCYTE CELL LINES AND RETINA.
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 CC DR EMBL; BC014423; AAA14423.1; -
 CC DR PIR; A55253; A55253
 CC DR Genew; HGNC; 7124; M1ANA.
 CC DR MIM; 605513; -
 CC DR GO; GO:0005887; C: integral to plasma membrane; TAS.
 CC DR GO; GO:0008222; F: tumor antigen; TAS.
 CC KW Antigen; Transmembrane.
 CC FT TRANSMEM 27
 CC SQ SEQUENCE 118 AA; 13157 MW; B755BF39CFB15E CRC64;
 CC
 CC Query Match 100.0%; Score 649; DB 1; Length 118;
 CC Best Local Similarity 100.0%; Pred. No. 3,4e-61;
 CC Matches 118; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 CC
 CC OY 1 MPRDAPFIYGYPRKGGHSTTREAAGIGILVYLLIGVLLICGWCRRNGRRLMDK 60
 CC ||||||||||||||||||||||||||||||||||||||||||||||||||||||||
 CC DB 1 MPRDAPFIYGYPRKGGHSTTREAAGIGILVYLLIGVLLICGWCRRNGRRLMDK 60
 CC ||||||||||||||||||||||||||||||||||||||||||||||||||||||||
 CC OY 61 SLHGTCALTRRCPOEGFDHDSKVSLOEKNCPEVYVNPAPAYEKLSAESPYPYSP 118
 CC ||||||||||||||||||||||||||||||||||||||||||||||||||||||||
 CC DB 61 SLHGTCALTRRCPOEGFDHDSKVSLOEKNCPEVYVNPAPAYEKLSAESPYPYSP 118
 CC ||||||||||||||||||||||||||||||||||||||||||||||||||||||||
 CC
 CC RESULT 2
 CC CD2_MOUSE STANDARD; PRT; 344 AA.
 CC AC P08920; 061394;
 CC DT 01-NOV-1988 (Rel. 09, Created)
 CC DT 01-NOV-1988 (Rel. 09, Last sequence update)
 CC DT 15-SEP-2003 (Rel. 42, Last annotation update)
 CC DE T-cell surface antigen CD2 precursor (T-cell surface antigen
 CC T11/Leu-5) (LFA-2) (LFA-3 receptor).
 CC GN CD2.
 CC OS Mus musculus (Mouse).
 CC OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
 CC OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.
 CC OX NCBI_TaxID:10090;
 CC RN [1]
 CC RP SEQUENCE FROM N.A.
 CC RC STRAIN-B10.A;
 CC RA MEDLINE-87276135; PubMed-2440689;
 CC RA Sewell W.A., Brown M.H., Fink P.J., Kozak C.A., Crumpton M.J.;
 CC "The murine homologue of the T lymphocyte CD2 antigen: molecular
 CC cloning, chromosome assignment and cell surface expression.";
 CC RT Eur. J. Immunol. 17:1015-1020(1987).
 CC RL [2]
 CC RN SEQUENCE FROM N.A.
 CC RP MEDLINE-88004738; PubMed-2820751;
 CC RA Clayton L.K., Sayre P.H., Novotny J., Reinherz E.L.;
 CC "Murine and human T11 (CD2) cDNA sequences suggest a common signal
 CC transduction mechanism.";
 CC RT Eur. J. Immunol. 17:1367-1370(1987).
 CC RL [3]
 CC RN SEQUENCE FROM N.A.
 CC RP STRAIN-BALB/c; TISSUE=Liver;
 CC RC MEDLINE-88144486; PubMed-2894031;
 CC RA Diamond D.J., Clayton L.K., Sayre P.H., Reinherz E.L.;
 CC "Ekon-inton organization and sequence comparison of human and murine
 CC T11 (CD2) genes.";
 CC RT Proc. Natl. Acad. Sci. U.S.A. 85:1615-1619(1988).
 CC RL

RN [4]
 RP SEQUENCE FROM N.A.
 RX MEDLINE-88140313; PubMed-3257775;
 RA Yagita H., Okumura K., Nakauchi H.;
 RT "Molecular cloning of the murine homologue of CD2. Homology of the
 RT molecule to its human counterpart T11.";
 RL J. Immunol. 140:1321-1326(1988).
 RN [5]
 RP INTERACTION WITH CD2AP.
 RX MEDLINE-98412662; PubMed-9741631;
 RA Wilder M.L., Olszowy M.W., Holdorf A.D., Li J., Bromley S., Desai N.,
 RA Dustin P., Rosenberger F., van der Merwe P.A., Allen P.M., Shaw A.S.;
 RT "A novel adaptor protein orchestrates receptor patterning and
 RT cytoskeletal polarity in T-cell contacts.";
 RL Cell 94:667-677(1998).
 CC -1- FUNCTION: CD2 interacts with lymphocyte function-associated
 CC antigen (LFA-3) and CD48/BCM1 to mediate adhesion between T
 CC cells and other cell types. CD2 is implicated in the triggering
 CC of T-cells, the cytoplasmic domain is implicated in the
 CC signaling function.
 CC -1- SUBUNIT: Interacts with CD2AP.
 CC -1- SUBCELLULAR LOCATION: Type I membrane protein.
 CC -1- SIMILARITY: Contains 1 immunoglobulin-like V-type domain.
 CC -1- SIMILARITY: Contains 1 immunoglobulin-like C2-type domain.
 CC -----
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 CC or send an email to license@isb-sib.ch).
 CC -----
 CC DR EMBL; Y00023; CAA68258.1; -
 CC DR EMBL; X06143; CAA29500.1; -
 CC DR EMBL; M19807; AAA37393.1; -
 CC DR EMBL; M19801; AAA37393.1; JOINED.
 CC DR EMBL; M19803; AAA37393.1; JOINED.
 CC DR EMBL; M19805; AAA37393.1; JOINED.
 CC DR EMBL; M18934; AAA37397.1; -
 CC DR PIR; 149585; 149585.
 CC DR HSSP; P08921; 1A64.
 CC DR GDI; MGI:88320; Cd2.
 CC DR GO; GO:0005515; F: protein binding activity; ISS.
 CC DR GO; GO:0004872; F: receptor activity; ISS.
 CC DR PROSITE; PS50835; IG-LIKE; FALSE_NDG.
 CC KW Immunoglobulin domain; T-cell; Glycoprotein; Antigen; Transmembrane;
 CC Cell adhesion; Repeat; Signal.
 CC FT SIGNAL 1 22
 CC FT CHAIN 23 344 T-CELL SURFACE ANTIGEN CD2.
 CC FT DOMAIN 23 203 EXTRACELLULAR (POTENTIAL).
 CC FT TRANSMEM 204 229 POTENTIAL.
 CC FT DOMAIN 230 344 CYTOPLASMIC (POTENTIAL).
 CC FT DOMAIN 23 121 IG-LIKE V-TYPE.
 CC FT DOMAIN 122 202 IG-LIKE C2-TYPE.
 CC FT DOMAIN 276 343 PRO-RICH.
 CC FT DISULFID 133 197 BY SIMILARITY.
 CC FT DISULFID 140 180
 CC FT CARBOHYD 82 82 N-LINKED (GLCNAC. . .) (POTENTIAL).
 CC FT CARBOHYD 94 94 N-LINKED (GLCNAC. . .) (POTENTIAL).
 CC FT CARBOHYD 135 135 N-LINKED (GLCNAC. . .) (POTENTIAL).
 CC FT CARBOHYD 166 166 N-LINKED (GLCNAC. . .) (POTENTIAL).
 CC FT CARBOHYD 99 99 Y -> T (IN REF. 3).
 CC FT CONFLICT 128 128 M -> V (IN REF. 3 AND 4).
 CC FT CONFLICT 139 139 T -> I (IN REF. 4).
 CC FT CONFLICT 175 175 N -> A (IN REF. 4).
 CC FT CONFLICT 175 175 N -> S (IN REF. 4).
 CC FT CONFLICT 191 191 K -> N (IN REF. 2).
 CC FT CONFLICT 192 192 M -> T (IN REF. 3 AND 4).
 CC SQ SEQUENCE 344 AA; 38414 MW; CFFD12FCHD1444450 CRC64;
 CC
 CC Query Match 12.4%; Score 80.5; DB 1; Length 344;

Best Local Similarity 27.4%; Pred. No. 0.44; Matches 31; Conservative 15; Mismatches 46; Indels 21; Gaps 5;

QY 3 REDAHFYGYKPKGHSHYTAEEAGIGITLVIGVLLICGWCRCRRNGYRALMDKSL 62
 Db 188 KESMEYVNCPEKLSF-YTVGVGAG-GLLIVL-VALFTFC-ICKRRKRRRRKDEL 243
 QY 63 HVGTCALTRRCPOEGFDRDSKVSIOEKNCPEVYPNAPPAEYKLSAEOSPP 115
 Db 244 EI-----KASRSTYVERGPKPHSTPAAANQNSVALQAPPP 279

RESULT 3

NASC_BACSU
 ID NASC_BACSU STANDARD; PRT; 710 AA.
 AC P42434;
 DT 01-NOV-1995 (Rel. 32, Created)
 DT 01-NOV-1997 (Rel. 35, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Asimilatory nitrate reductase catalytic subunit (EC 1.7.99.4).
 GN NASC OR NARB OR NASBB.
 OS Bacillus subtilis.
 OC Bacteria; Firmicutes; Bacillales; Bacillaceae; Bacillus.
 OX NCBI_TaxID=1423;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=168;
 RX MEDLINE=97124189; PubMed=8969502;
 RA Yamane K., Kumano M., Kurita K.;
 RT "The 25 degrees-36 degrees region of the Bacillus subtilis
 RT chromosome: determination of the sequence of a 146 kb segment and
 RT identification of 113 genes.";
 RL Microbiology 142:3047-3056(1996).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN=168;
 RX MEDLINE=98044033; PubMed=9384377;
 RA Kunst F., Ogasawara N., Moszer I., Albertini A.M., Alloni G.,
 RA Azevedo V., Bertero M.G., Bessieres P., Bolotin A., Borchert S.,
 RA Boriss R., Boursier L., Brans A., Braun M., Brigelli S.C., Bron S.,
 RA Brouillet S., Bruschi C.V., Caldwell B., Capuano V., Carter N.M.,
 RA Choi S.K., Codani J.J., Connerton I.F., Cummings N.J., Daniel R.A.,
 RA Deniot F., Devigne K.M., Dusterhoft A., Ehrlich S.D., Emerson P.T.,
 RA Ertlan K.D., Errington J., Fabret C., Ferrari E., Foulger D.,
 RA Frits C., Fujita M., Fujita Y., Fuma S., Galizzi A., Galleron N.,
 RA Ghim S.Y., Glaser P., Goffeau A., Golightly E.J., Grandi G.,
 RA Giuseppe G., Guy B.J., Haga K., Halech J., Harwood C.R., Henaut A.,
 RA Hilbert H., Holsappel S., Hosono S., Hullo M.F., Itaya M., Jones L.,
 RA Joris B., Katsuma D., Kasahara Y., Klastri-Blanchard M., Klein C.,
 RA Kobayashi Y., Koetter P., Koningsstein G., Krogh S., Kumano M.,
 RA Kurita K., Lapidus A., Lardinois S., Lauber J., Lazarevic V.,
 RA Lee S.M., Levine A., Liu H., Masuda S., Manuel C., Medigue C.,
 RA Medina N., Mellado R.P., Mizuno M., Moesti D., Nakai S., Nodack M.,
 RA Noone D., O'Reilly M., Ogawa K., Ogiwara A., Oudega B., Park S.H.,
 RA Paro V., Pohl T.M., Portetelle D., Porwollik S., Prescott A.M.,
 RA Prescan E., Pujic P., Purnelle B., Rapoport G., Rey M., Reynolds S.,
 RA Rieger M., Rivolta C., Rocha E., Roche B., Rose M., Sadleir Y.,
 RA Sato T., Scanlan E., Schleich S., Schroeter R., Scofield F.,
 RA Sekiguchi J., Sekowska A., Seror S.J., Serron P., Shin B.S., Solido B.,
 RA Sorokin A., Tacconi E., Takagi T., Takahashi H., Takemaru K.,
 RA Takeuchi M., Tamakoshi A., Tanaka T., Terpstra P., Tognoni A.,
 RA Tosato V., Uchiyama S., Vandenbol M., Vanlier F., Vassartoli A.,
 RA Viari A., Wambutt R., Wedler H., Wedler F., Weitzenecker T.,
 RA Winters P., Wipat A., Yamamoto H., Yamane K., Yasumoto K., Yata K.,
 RA Yoshida K., Yoshikawa H.F., Zumslein E., Yoshikawa H., Danchin A.,
 RT "The complete genome sequence of the Gram-positive bacterium Bacillus
 RT subtilis.";
 RL Nature 390:249-256(1997).
 RN [3]
 RP SEQUENCE OF 35-710 FROM N.A.
 RC STRAIN=168;
 RX MEDLINE=95173124; PubMed=7866621;
 RA Ogawa K.-I., Akagawa E., Yamane K., Sun Z.-W., Iacelle M., Zuber P.,

RA Nakano M.M.;
 RT "The nasB operon and nasB gene are required for nitrate/nitrite
 RT assimilation in Bacillus subtilis.";
 RL J. Bacteriol. 177:1409-1413(1995).
 CC -1- FUNCTION: NITRATE REDUCTASE IS A KEY ENZYME INVOLVED IN THE FIRST
 CC STEP OF NITRATE ASSIMILATION IN PLANTS, FUNGI AND BACTERIA.
 CC -1- CATALYTIC ACTIVITY: Nitrite + acceptor -> nitrate + reduced
 CC acceptor.
 CC -1- COFACTOR: MOLYBDENUM (MOLYBDOPTERIN); MAY BIND A 4FE-4S CLUSTER.
 CC -1- PATHWAY: Nitrate assimilation (identification).
 CC -1- SIMILARITY: BELONGS TO THE PROKARYOTIC MOLYBDOPTERIN-CONTAINING
 CC OXIDOREDUCTASE FAMILY.
 CC -----
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 CC -----
 CC DR EMBL: D50453; BAA08965.1; -
 CC DR EMBL: Z99105; CAB12125.1; -
 CC DR EMBL: D30689; BAA06353.1; -
 CC DR PIR: E69665; E69665.
 CC DR HSSP: P81186; 2NAP.
 CC DR Subtilist; BG11095; nasC.
 CC DR InterPro; IPR006657; Mol_dinuc_bind.
 CC DR InterPro; IPR006963; Molybdop_Fe4S4.
 CC DR InterPro; IPR006656; Molybdopterin.
 CC DR InterPro; IPR006655; Prok_Moxred.
 CC DR Pfam; PF04879; Molybdop_Fe4S4; 1.
 CC DR Pfam; PF00384; molybdopterin; 1.
 CC DR Pfam; PF01568; Molybdop_binding; 1.
 CC DR PROSITE; PS00551; MOLYBDOPTERIN_PROK_1; 1.
 CC DR PROSITE; PS00490; MOLYBDOPTERIN_PROK_2; 1.
 CC DR PROSITE; PS00932; MOLYBDOPTERIN_PROK_3; FALSE NEG.
 CC DR Oxidoreductase; Molybdenum; Nitrate assimilation; Iron-sulfur; 4Fe-4S;
 CC KW Complete proteome.
 CC FT METAL 26 26 IRON-SULFUR (4FE-4S) (POTENTIAL).
 CC FT METAL 29 29 IRON-SULFUR (4FE-4S) (POTENTIAL).
 CC FT METAL 33 33 IRON-SULFUR (4FE-4S) (POTENTIAL).
 CC FT METAL 63 63 IRON-SULFUR (4FE-4S) (POTENTIAL).
 CC SQ SEQUENCE 710 AA; 78575 MW; 625E8664A1552AA2 CRC64;
 CC -----
 CC Query Match 11.9%; Score 77; DB 1; Length 710;
 CC Best Local Similarity 32.4%; Pred. No. 2.2;
 CC Matches 34; Conservative 8; Mismatches 37; Indels 26; Gaps 6;
 QY 15 KGHGSHYTAEE-----AAGIGITLVIGVLLIGCWT--CRRNGYRALMDKSLNVT 66
 Db 507 KGRFYTSADIDFENELREASRG-----GIADYSGISYGRLRGGIHMPCPESDHPET 560
 QY 67 QCATLRRCPOEGFDRDSKVSIOEKNCPEVYPNAPPA-YEKLSAE 110
 Db 561 GRFLT-----ESFNPDPKALNS-----VIPNEPVRKEKPTAD 594
 RESULT 4
 AD19_HUMAN
 ID AD19_HUMAN STANDARD; PRT; 956 AA.
 AC Q9H013; Q9BZL5; Q9UHP2;
 DT 28-FEB-2003 (Rel. 41, Created)
 DT 28-FEB-2003 (Rel. 41, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE ADAM 19 precursor (BC 3.4.24.-) (A disintegrin and metalloproteinase
 DE domain 19) (Mellitin beta) (Metalloprotease and disintegrin dentritic
 DE antigen marker) (MADDAM).
 GN ADAM19 OR MLTNB OR PRSG34.
 OS Homo sapiens (Human).
 CC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 CC Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
 OX NCBI_TaxID=9606;

[illegible]

ID AD15_MOUSE STANDARD; PRT; 815 AA.
 AC 088839; 090YL2;
 DT 16-OCT-2001 (Rel. 40, Created)
 DT 16-OCT-2001 (Rel. 40, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE ADAM 15 precursor (EC 3.4.24.-) (A disintegrin and metalloproteinase domain 15) (Metalloproteinase-like, disintegrin-like, and cysteine-rich protein 15) (MDC-15) (Metalloproteinase RGD disintegrin protein) (Metargidin) (AD56).
 DE ADAM15 OR MDC15.
 GN Mus musculus (Mouse).
 OS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 OC NCBI_TaxID:10090;
 RN [1]
 RP SEQUENCE FROM N.A., AND PROCESSING.
 RC TISSUE=Lung;
 RX MEDLINE=98421554; PubMed=9748307;
 RA Lum L., Reid M.S., Blobel C.P.;
 RT "Intracellular maturation of the mouse metalloproteinase disintegrin MDC15.";
 RT J. Biol. Chem. 273:26236-26247 (1998).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Myeloid, and Myeloma;
 RA Shimizu E., Higuchi Y., Matsuura K., Hijiya N., Yamamoto S.;
 RT "Structure of the mouse ADAM 15 (AD56) gene.";
 RT Submitted (JAN-1999) to the EMBL/GenBank/DBJ databases.
 RN [3]
 RP INTERACTION WITH ENDOPHILIN I AND SNX9.
 RX MEDLINE=20002705; PubMed=10531379;
 RA Howard L., Nelson K.K., Maciewicz R.A., Blobel C.P.;
 RT "Interaction of the metalloproteinase disintegrins MDC9 and MDC15 with two SH3 domain-containing proteins, endophilin I and SHPXL.";
 RT J. Biol. Chem. 274:31693-31699 (1999).
 CC -1- FUNCTION: May be involved in cell-surface proteolysis, cell adhesion or intracellular protein maturation.
 CC -1- COFACTOR: Binds 1 zinc ion per subunit (Potential).
 CC -1- SUBUNIT: Interacts specifically with Src family protein-tyrosine kinases (PTKs) (By similarity). Interacts with ITAG-ITGB3 (Vitronection receptor), PACSIN3 and SNX9. PACSIN3 and SNX9 preferentially bind the precursor but not the processed form of ADAM15, suggesting that the interaction occurs in a secretory pathway compartment prior to the medial Golgi.
 CC -1- SUBCELLULAR LOCATION: TYPE I MEMBRANE PROTEIN. THE MAJORITY OF THE PROTEIN IS LOCALIZED IN A PERINUCLEAR COMPARTMENT WHICH MAY CORRESPOND TO THE TRANS-GOLGI NETWORK OR THE LATE ENDOSOME. THE PRO-PROTEIN IS THE MAJOR DETECTABLE FORM ON THE CELL SURFACE, WHEREAS THE MAJORITY OF THE PROTEIN IN THE CELL IS PROCESSED.
 CC -1- TISSUE SPECIFICITY: HIGHLY EXPRESSED IN HEART, BRAIN, LUNG, AND KIDNEY. EXPRESSED AT LOWER LEVELS IN SPLEEN, LIVER, TESTIS AND MUSCLE.
 CC -1- DOMAIN: THE CYTOPLASMIC DOMAIN INTERACTS WITH ENDOPHILIN I AND SORTING NEXIN 9.
 CC -1- DOMAIN: DESINTEGRIN DOMAIN BINDS TO INTEGRIN ALPHA-V-BETA3 (BY SIMILARITY).
 CC -1- PTM: THE PRECURSOR IS CLEAVED BY A FURIN ENDOPEPTIDASE. AN ADDITIONAL MEMBRANE PROXIMAL SITE OF CLEAVAGE AFFECTS A SMALL PERCENTAGE OF THE PROTEIN AND RESULTS IN DISULFIDE-LINKED FRAGMENTS. THE PRO-DOMAIN IS APPARENTLY CLEAVED IN SEVERAL POSITIONS THAT ARE N-TERMINAL OF THE FURIN CLEAVAGE SITE.
 CC -1- PTM: MAY BE PARTIALLY SIALYLATED.
 CC -1- PTM: Phosphorylation increases association with PTKs (By similarity).
 CC -1- SIMILARITY: Belongs to peptidase family M12B.
 CC -1- SIMILARITY: Contains 1 EGF-like domain.
 CC -1- SIMILARITY: Contains 1 disintegrin domain.
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 CC -----
 CC EMBL: AF006196; AAC61896.1; -;
 CC EMBL: AB022089; BAA86903.1; -;
 CC HSSP: P17494; 1KST.
 CC MEROPS: M12.215; -;
 CC MGD: MGI:1333882; Adam15.
 CC InterPro: IP0006586; ADAM_cysteine.
 CC InterPro: IP0001762; Disintegrin.
 CC InterPro: IP0006209; EGF-like.
 CC InterPro: IP0001818; Matrixin.
 CC InterPro: IP0002870; Pep_M12B_propep.
 CC InterPro: IP0001590; Repolysin.
 CC InterPro: IP0006025; zn_MTPeptide.
 CC Pfam: PF00200; disintegrin; 1.
 CC Pfam: PF01562; Pep_M12B_propep; 1.
 CC Pfam: PF01421; Repolysin; 1.
 CC ProDom: PD000664; Disintegrin; 1.
 CC SMART: SM00608; ACR; 1.
 CC SMART: SM00050; DISIN; 1.
 CC PROSITE: PS50215; ADAM_MEPPO; 1.
 CC PROSITE: PS00022; EGF_1; FALSE_NEG.
 CC PROSITE: PS01186; EGF_2; 1.
 CC PROSITE: PS00427; DISINTEGRIN_1; FALSE_NEG.
 CC PROSITE: PS50214; DISINTEGRIN_2; 1.
 CC PROSITE: PS00142; ZINC_PROTEASE; 1.
 CC PROSITE: PS00546; CYSTEINE_SWITCH; FALSE_NEG.
 CC Hydrolase; Metalloproteinase; zinc; Signal; Glycoprotein; Zymogen;
 CC Transmembrane; EGF-like domain; SH3-binding; Phosphorylation.
 CC SIGNAL 1 17
 CC PROPEP 18 207
 CC CHAIN 18 815
 CC DOMAIN 208 696
 CC TRANSMEM 697 717
 CC DOMAIN 718 815
 CC DOMAIN 422 509
 CC DOMAIN 510 657
 CC DOMAIN 658 686
 CC DOMAIN 689 712
 CC SITE 767 773
 CC SITE 802 808
 CC SITE 179 179
 CC SITE 289 290
 CC METAL 349 349
 CC METAL 350 350
 CC ACT_SITE 350 353
 CC METAL 353 359
 CC METAL 324 410
 CC DISULFID 481 494
 CC DISULFID 658 668
 CC DISULFID 662 674
 CC DISULFID 676 685
 CC MOD_RES 716 716
 CC MOD_RES 736 736
 CC CARBOHYD 238 238
 CC CARBOHYD 390 390
 CC CARBOHYD 393 393
 CC CARBOHYD 607 607
 CC CARBOHYD 612 612
 CC CONFLICT 21 22
 CC CONFLICT 443 443
 CC CONFLICT 459 459
 CC CONFLICT 564 565
 CC CONFLICT 654 660
 CC CONFLICT 660 660
 CC CONFLICT 703 703
 CC CONFLICT 712 712
 CC CONFLICT 729 729
 CC CONFLICT 797 797
 CC CONFLICT 803 805
 CC CONFLICT 810 810
 CC ADAM 15.
 CC EXTRACELLULAR (POTENTIAL).
 CC POTENTIAL.
 CC CYTOPLASMIC (POTENTIAL).
 CC METALLOPROTEASE.
 CC DISINTEGRIN-LIKE.
 CC CYS-RICH.
 CC EGF-LIKE.
 CC POLY-LEU.
 CC SH3-BINDING (POTENTIAL).
 CC SH3-BINDING (POTENTIAL).
 CC CYSTEINE SWITCH.
 CC CLEAVAGE (BY PERIN) (POTENTIAL).
 CC ZINC (CATALYTIC) (POTENTIAL).
 CC POTENTIAL.
 CC ZINC (CATALYTIC) (POTENTIAL).
 CC ZINC (CATALYTIC) (POTENTIAL).
 CC BY SIMILARITY.
 CC BY SIMILARITY.
 CC BY SIMILARITY.
 CC PHOSPHORYLATION (BY SIMILARITY).
 CC N-LINKED (GLYCANC. . .) (POTENTIAL).
 CC N-LINKED (GLYCANC. . .) (POTENTIAL).
 CC N-LINKED (GLYCANC. . .) (POTENTIAL).
 CC N-LINKED (GLYCANC. . .) (POTENTIAL).
 CC N-LINKED (GLYCANC. . .) (POTENTIAL).
 CC N-LINKED (GLYCANC. . .) (POTENTIAL).
 CC PP -> RR (IN REF. 2).
 CC E -> Q (IN REF. 2).
 CC G -> E (IN REF. 2).
 CC SP -> T (IN REF. 2).
 CC G -> E (IN REF. 2).
 CC R -> S (IN REF. 2).
 CC L -> R (IN REF. 2).
 CC L -> R (IN REF. 2).
 CC L -> S (IN REF. 2).
 CC PAP -> AAS (IN REF. 2).
 CC A -> P (IN REF. 2).

SQ SEQUENCE 815 AA; 87424 MW; C064BD3B7347D19B CRC64;
 Query Match 11.2%; Score 72.5; DB 1; Length 815;
 Best Local Similarity 20.9%; Pred. No. 7.7;
 Matches 28; Conservative 17; Mismatches 40; Indels 49; Gaps 5;

QY 14 KKGHSHY-----TTAEAGIGILTVILGVLILGICWYCR 50
 DB 660 RKCHGVCCSSGRCBEGNAPDCMTQKATSSLTGILLVLLASTYHR 719
 QY 51 RRGYRALMDKSLHVTGOCALTRRCPOBGFDRHDSKVSLSQEKNCBPVPNPAPAYEKLSAE 110
 DB 720 ARLEHRLCQLK---GSSCQY---RAPQ-----SCPPERGPFOBAQOMGT 759
 QY 111 QS-----PPPYSP 118
 DB 760 KSGGPRKPPPPRP 773

RESULT 6
 EXPL_DROME STANDARD; PRT: 1429 AA.
 AC 007436;
 DT 01-OCT-1996 (Rel. 34, Created)
 DT 01-NOV-1997 (Rel. 35, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Expanded protein.
 GN Ex.
 OS Drosophila melanogaster (Fruit fly).
 OC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
 OC Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
 OC Ephydroidea; Drosophilidae; Drosophila.
 NCBI_TaxID=7227;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC TISSUE-Imaginal disks;
 RX MEDLINE=94094747; PubMed=8269855;
 RA Boedihelmer M., Laugheon A.;
 RT "Expanded: a gene involved in the control of cell proliferation in
 RL Imaginal discs.";
 RU Development 118:1291-1301(1993).
 RN [2]
 RP REVISIONS.
 RA Boedihelmer M.;
 RT Submitted (DEC-1996) to the EMBL/GenBank/DBJ databases.
 CC -1- FUNCTION: INVOLVED IN THE CONTROL OF CELL PROLIFERATION IN
 CC IMAGINAL DISCS. MAY BIND TO CERTAIN PROTEINS OF SIGNAL
 CC TRANSDUCTION PATHWAYS BY INTERACTION WITH THEIR SH3 DOMAINS.
 CC -1- SUBCELLULAR LOCATION: APICAL SURFACE OF DISC CELLS.
 CC -1- DISEASE: MUTATIONS OF EXPANDED PROTEIN CAUSE HYPERPLASIA OF THE
 CC IMAGINAL DISC RESULTING IN WING OVERGROWTH. THIS OVERGROWTH IS
 CC LIMITED TO SPECIFIC REGIONS ALONG THE 2 WING AXES. DEFECTS ALSO
 CC IN EYES, HEAD, THORAX AND LIMBS WHERE DUPLICATION AND BULGING
 CC OFTEN OCCUR.
 CC -1- SIMILARITY: Contains 1 FERM domain.
 CC -----
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 CC or send an email to license@isb-sib.ch).
 CC -----
 CC EMBL: L14768; AAB3974.1; -
 CC DR F13720: 113720.
 CC DR FlyBase: FBgn0004583; ex.
 CC DR InterPro: IPR000299; Band.4.1.
 CC DR Pfam: PF00373; Band.41; 1.
 CC DR SMART: SM00295; B41; 1.
 CC DR PROSITE: PS00660; FERM_1; FALSE_NEG.
 CC DR PROSITE: PS00661; FERM_2; FALSE_NEG.
 CC DR PROSITE: PS50057; FERM_3; 1.

KM Developmental protein; SH3-binding.
 FT DOMAIN 26 399 FERM.
 FT SITE 1008 1016 SH3-BINDING (POTENTIAL).
 FT SITE 1012 1020 SH3-BINDING (POTENTIAL).
 FT SITE 1149 1157 SH3-BINDING (POTENTIAL).
 FT DOMAIN 409 412 POLY-GLU.
 FT DOMAIN 782 788 POLY-PRO.
 FT DOMAIN 952 955 POLY-HIS.
 FT DOMAIN 1002 1005 POLY-PRO.
 FT DOMAIN 1011 1017 POLY-PRO.
 FT DOMAIN 1081 1084 POLY-PRO.
 FT DOMAIN 1149 1154 POLY-PRO.
 FT DOMAIN 1158 1168 POLY-ALA.
 FT DOMAIN 1170 1174 POLY-SER.
 FT DOMAIN 1199 1205 POLY-PRO.
 FT DOMAIN 1416 1424 POLY-GLN.
 SQ SEQUENCE 1429 AA; 153886 MW; 3CB08D2FC4862062 CRC64;

Query Match 11.2%; Score 72.5; DB 1; Length 1429;
 Best Local Similarity 23.4%; Pred. No. 14;
 Matches 37; Conservative 10; Mismatches 52; Indels 59; Gaps 7;

QY 2 PREDARFTYGPYPRKGGH-----HSYTAERAGIGILTVILGVLILGICWYCR 54
 DB 708 PRSDNVSTGSSFRDGDSPDTNKHSLSAEFLTN-----LIVG-----RGTY 750
 QY 55 --RALMDKSLH-----VTGOCALTRRCPOBGFDRHDSKVSLSL----- 88
 DB 751 PSRKTYSLSLHSDCDYVTLPLDQGEEDVDPAPPAPPPYSAHEKTYGLGCPPIAKPIPKP 810
 QY 89 -----QEKNCBPVPNPAP-PAYEKLSAEOSPPYYS 117
 DB 811 IAVVAPKPDSPPCSPVPVPAIPAPPAIRRRDPPYYS 848

RESULT 7
 PCH7_HUMAN STANDARD; PRT: 1069 AA.
 AC 060245; 060245; 060247;
 DT 28-FEB-2003 (Rel. 41, Created)
 DT 28-FEB-2003 (Rel. 41, Last sequence update)
 DT 15-SEP-2003 (Rel. 42, Last annotation update)
 DE Protocadherin 7 precursor (Brain-heart protocadherin) (BH-Pcdh).
 GN PCDH7 OR BHPCDH.
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A. (ISOFORMS A; B AND C).
 RX MEDLINE=98277460; PubMed=9615233;
 RA Yoshida K., Yoshitomo-Nakagawa K., Seki N., Sasaki M., Sugano S.;
 RT "Cloning, expression analysis, and chromosomal localization of
 RT BH-protocadherin (PCDH7), a novel member of the cadherin
 RT superfamily.";
 RT Genomics 49:458-461(1998).
 RL Genomics 49:458-461(1998).
 CC -1- SUBCELLULAR LOCATION: Type I membrane protein.
 CC -1- ALTERNATIVE PRODUCTS:
 CC Event-Alternative splicing; Named isoforms-3;
 CC Name-A: Synonyms-BH-Pcdh-a;
 CC IsoId=060245-1; Sequence-Displayed;
 CC Name-B: Synonyms-BH-Pcdh-b;
 CC IsoId=060245-2; Sequence-VSP_000704;
 CC Name-C: Synonyms-BH-Pcdh-c; Sequence-VSP_000705; VSP_000706;
 CC IsoId=060245-3; Sequence-VSP_000706;
 CC -1- TISSUE SPECIFICITY: Expressed predominantly in brain and heart and
 CC at lower levels in various other tissues.
 CC -1- SIMILARITY: Contains 7 cadherin domains.
 CC -----
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DT 28-FEB-2003 (Rel. 41, Created)
 DT 28-FEB-2003 (Rel. 41, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Transmembrane gamma-carboxylglutamic acid protein 1 precursor (Proline-rich
 DE rich Glu protein 1) (Proline-rich gamma-carboxylglutamic acid protein
 DE 1)
 GN PRK1 OR TWG1 OR PRP1.
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 OX NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA MEDLINE-97404347; PubMed-9256434;
 RA Kulman J.D., Harris J.E., Haldeman B.A., Dave E.W.;
 RT "Primary structure and tissue distribution of two novel proline-rich
 RT gamma-carboxylglutamic acid proteins."
 RT Proc. Natl. Acad. Sci. U.S.A. 94:9058-9062(1997).
 CC -1- PTM: Glu residues are produced after subsequent posttranslational
 CC modifications of glutamic acid by a vitamin K-dependent gamma-
 CC carboxylase.
 CC -----
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 CC or send an email to license@isb-sib.ch).
 CC -----
 CC EMBL: AF009242; AAB67070.1; -
 CC DR HSSP: P00740; ICFH.
 CC DR GeneW: HGNC:9469; PRK1.
 CC DR MIM: 604428; -
 CC GO: GO:0005887; C: integral to plasma membrane; TAS.
 CC InterPro: IPR002383; GLA_blood.
 CC InterPro: IPR000294; Vitk_dep_GLA.
 CC Pfam: PF00594; gla; 1.
 CC PRINTS: PR00001; GLABLOOD.
 CC SMART: SM00069; GLA; 1.
 CC DR PROSITE: PS00011; GLT_CARBOXYLATION; 1.
 CC DR Gamma-carboxylglutamic acid; Vitamin K; Transmembrane.
 CC FT PROPEP 1 20
 CC FT CHAIN 1 218
 CC FT TRANSMEMBRANE GAMMA-CARBOXYGLUTAMIC ACID
 CC FT PROTEIN 1.
 CC FT DOMAIN 21 83 EXTRACELLULAR (POTENTIAL).
 CC FT TRANSMEM 84 106 POTENTIAL.
 CC FT DOMAIN 107 218 CYTOPLASMIC (POTENTIAL).
 CC FT DOMAIN 24 61 GLA-RICH.
 CC FT DOMAIN 131 135 POLY-PRO.
 CC FT SEQUENCE 218 AA; 24947 MW; 26538A61A80AEB98 CRC64;
 SQ
 Query Match 10.7%; Score 69.5; DB 1; Length 218;
 Best Local Similarity 27.0%; Pred. No. 3.9;
 Matches 30; Conservative 9; Mismatches 37; Indels 35; Gaps 5;
 Oy 33 LTVIIGVLLIGCWCRNGRYALMDKSLHWGTCALTRRCPOEGFDH-----PP 81
 Db 91 LFTIILVYILT--WRCFILANKTRRQIVTSGHIPFPHALNITPPPPDEYDSSGSLSPGR 148
 Oy 82 -----RDSKVSILQENKCEPVVNPAPAYEKLSAEGS-----PPP 115
 Db 149 LQYVGRSDSVSTRLSNCDP-----PPTYEATGQVNLQRSETEPHILPPP 194
 RESULT 10
 LSGL_HAEIN STANDARD: PRT; 401 AA.
 AC P71399; Q48210;
 ID 16-OCT-2001 (Rel. 40, Created)
 DT 16-OCT-2001 (Rel. 40, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)

DE Isg locus putative protein 1.
 GN H11700.
 OS Haemophilus influenzae.
 OC Bacteria; Proteobacteria; Gammaproteobacteria; Pasteurellales;
 OC Pasteurellaceae; Haemophilus.
 OX NCBI_TaxID=727;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA STRAIN-A2;
 RC McLaughlin R., Abu Kwaik Y., Young R., Splinola S., Apicella M.;
 RT "Characterization and sequence of the Isg locus from Haemophilus
 RT influenzae."
 RT Submitted (JUN-1992) to the EMBL/GenBank/DBJ databases.
 RN [2]
 RP SEQUENCE FROM N.A.
 RA STRAIN-Rd / KW20 / ATCC 51907;
 RC MEDLINE-95305630; PubMed-7542800;
 RA Fleischmann R.D., Adams M.D., White O., Clayton R.A., Kirkness E.F.,
 RA Kervage A.R., Bult C.J., Tomb J.-F., Dougherty B.A., Merrick J.M.,
 RA McKenney K., Sutton G., Fitzhugh W., Fields C.A., Gocayne J.D.,
 RA Scott J.D., Shirley R., Liu L.-I., Glodek A., Kelley J.M.,
 RA Weidman J.F., Phillips C.A., Spriggs T., Hedblom E., Cotton M.D.,
 RA Uterback T.R., Hanna M.C., Nguyen D.T., Sadek D.M., Brandon R.C.,
 RA Fine L.D., Fritchman J.L., Fritchman J.L., Geoghegan N.S.M.,
 RA Gehler C.L., McDonald L.A., Small K.V., Fraser C.M., Smith H.O.,
 RA Venter J.C.;
 RA "Whole-genome random sequencing and assembly of Haemophilus influenzae
 RA Rd."
 RT Science 269:496-512(1995).
 CC -1- SUBCELLULAR LOCATION: Integral membrane protein (Potential).
 CC -1- SIMILARITY: Belongs to the polysaccharide synthase family.
 CC H10867/H11700 SUBFAMILY.
 CC -----
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 CC -----
 CC EMBL: M94855; AAA24978.1; -
 CC DR EMBL: U32842; AAC23346.1; ALT_INTT.
 CC DR PIR: H64175; H64175.
 CC DR TIGR: H11700; -
 CC DR InterPro: IPR002797; Polysacc_synt.
 CC DR Pfam: PF01943; Polysacc_synt; 1.
 CC KW Hypothetical protein, Transmembrane. Complete proteome.
 CC FT TRANSMEM 8 28
 CC FT TRANSMEM 36 56
 CC FT TRANSMEM 87 107
 CC FT TRANSMEM 132 152
 CC FT TRANSMEM 162 182
 CC FT TRANSMEM 199 219
 CC FT TRANSMEM 237 257
 CC FT TRANSMEM 282 302
 CC FT TRANSMEM 320 340
 CC FT TRANSMEM 352 372
 CC FT TRANSMEM 374 394
 CC FT TRANSMEM 249 269
 CC FT TRANSMEM 276 296
 CC FT TRANSMEM 358 378
 CC FT TRANSMEM 401 AA; 45944 MW; FE2E7B02747B0874 CRC64;
 SQ
 Query Match 10.7%; Score 69.5; DB 1; Length 401;
 Best Local Similarity 21.6%; Pred. No. 7.5;
 Matches 21; Conservative 20; Mismatches 23; Indels 33; Gaps 4;
 Oy 6 AHRYGPKR-----GHGSHYTAEEAAGIGITVLIGVLLIGCW-----YCRRR 51
 Db 63 SRFFYFGKRSIMNVYVGTAYIT-----IIGSTIILGICWIAOSEILFYALNS 110
 Oy 52 NGYRALMDKSLHWGTCALTRRCPOEGFDRHDSKVS 88

Db 111 SIFOSFLNVLSTV-----RQCCKKMSYAFIOFSL 140

RESULT 11
CYTL_MOUSE STANDARD: PRT; 165 AA.

AC 08VHT7;
DT 28-FEB-2003 (Rel. 41, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Cysteine and tyrosine-rich protein 1 precursor (Proline-rich domain containing protein).
DE CYR1.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_Taxid=10090;
[1]

RP SEQUENCE FROM N.A.
RC STRAIN-BALB/C; TISSUE-Brain;
RX MEDLINE=22057895; PubMed=12062809;
RA Vitale L., Casadei R., Canaider S., Lenzi L., Strippoli P.,
D'Addabbo P., Glanone S., Carinci P., Zanolli M.;
RT "Cysteine and tyrosine-rich 1 (CYR1), a novel unpredicted gene on human chromosome 21 (21q21.2), encodes a cysteine and tyrosine-rich protein and defines a new family of highly conserved vertebrate specific genes.";
RL Gene 290:141-151(2002).
[2]

RP SEQUENCE FROM N.A.
RX MEDLINE=22032984; PubMed=12036297;
RA Raymond A., Camargo A.A., Deutsch S., Stevenson B.J., Parmigiani R.B.,
Ucla C., Bettont F., Kossler C., Lyle R., Guipont M., de Souza S.,
RA Iseli C., Jongeneel C.V., Bucher P., Simpson A.J.G.,
RA Antonarakis S.E.;
RT "Nineteen additional unpredicted transcripts from human chromosome 21.";
RL Genomics 79:824-832(2002).

CC -I- SUBCELLULAR LOCATION: Type I membrane protein (Potential).
CC -----

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CC -----

CC EMBL: AF442733; AAL35295.1;
DR EMBL: AY061854; AAL35738.1;
DR MGD: MGI:2152187; Cyrl.
KW Signal; Transmembrane.

FT SIGNAL 1 29 POTENTIAL.
FT CHAIN 30 165 CYSTEINE AND TYROSINE-RICH PROTEIN 1.
FT DOMAIN 30 61 EXTRACELLULAR (POTENTIAL).
FT TRANSMEM 62 82 POTENTIAL.
FT DOMAIN 83 165 CYTOPLASMIC (POTENTIAL).
FT DOMAIN 144 149 POLY-PRO.
FT CONFLICT 36 36 R -> H (IN REF. 2).
SO SEQUENCE 165 AA; 18063 MM; 94940DAAEZBCFDCI CRC64;

Query Match 10.5%; Score 68; DB 1; Length 165;
Best Local Similarity 21.2%; Pred. No. 4.2;
Matches 22; Conservative 19; Mismatches 37; Indels 26; Gaps 5;

QY 28 AGIGILVITGVLILG-----CWYCRRRNGYALMDKSLHVGQCALTRCPQEG 78
DB 59 SGRALAGIVGIVFIMVINGIALICICMKNRGTGIVRAAHINA-ISTPMAPPYT 117
QY 79 FDRH-DSKVSIOEKNCPEVVPNAPAYE---KLSAESPPPYSP 118
DB 118 YDHMEYRTDL-----PPYSAPQASAGSSPPPY 149

RESULT 12
COME_METUA STANDARD: PRT; 188 AA.

AC P58416; Q57704;
DT 28-FEB-2003 (Rel. 41, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Sulfolysate decarboxylase subunit beta (EC 4.1.1.79).
GN COME OR MJ0256.
OS Methanococcus jannaschli.
OC Archaea; Euryarchaeota; Methanococci; Methanococcales;
OC Methanocaldococcaceae; Methanocaldococcus.
OX NCBI_Taxid=2190;
[1]

RP SEQUENCE FROM N.A.
RC STRAIN-JAL-1 / DSM 2661 / ATCC 43067;
RX MEDLINE=96337999; PubMed=8688087;
RA Bult C.J., White O., Olsen G.J., Zhou L., Fleischmann R.D.,
Sutton G.G., Blake J.A., Fitzgerald L.M., Clayton R.A., Gocayne J.D.,
Kerlavage A.R., Dougherty B.A., Tomb J.F., Adams M.D., Reich C.I.,
Overbeek R., Kirkness E.F., Weinstock K.G., Merrick J.M., Glodek A.,
RA Scott J.L., Geoghagen N.S.M., Weidman J.F., Fuhrmann J.L., Nguyen D.,
Utterback T.R., Kelley J.M., Peterson J.D., Sadow P.W., Hanna M.C.,
RA Cotton M.D., Roberts K.M., Hirst M.A., Kaine B.P., Borodovsky M.,
Klenk H.-P., Fraser C.M., Smith H.O., Woese C.R., Venter J.C.;
RT "Complete genome sequence of the methanogenic archaeon, Methanococcus jannaschli.";
RL Science 273:1058-1073(1996).
[2]

RP CHARACTERIZATION.
RX MEDLINE=20398170; PubMed=10940029;
RA Graupner M., Xu H., White R.H.;
RT "Identification of the gene encoding sulfolysate decarboxylase, an enzyme involved in biosynthesis of coenzyme M.";
RL J. Bacteriol. 182:4862-4867(2000).

CC -I- FUNCTION: Catalyzes the decarboxylation of sulfolysate acid to sulfoacetalddehyde.
CC -I- CATALYTIC ACTIVITY: 3-sulfolysate - 2-sulfoacetalddehyde + CO(2).
CC -I- COFACTOR: TPP (Potential).
CC -I- PATHWAY: Coenzyme M biosynthesis; fourth step.
CC -I- SUBUNIT: Heterododecamer composed of 6 subunits alpha and 6 subunits beta.
CC -I- MISCELLANEOUS: Inactivated by oxygen when heated in air at 80 degrees Celsius.
CC -I- SIMILARITY: BELONGS TO THE TPP ENZYME FAMILY.
CC -I- CAUTION: The sequence corresponding to this entry was originally deleted in July 1999 because TIGR removed the CDS for that ORF. We have recreated it because of the evidence (ref.2) that it really exists.
CC -----

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CC -----

CC EMBL: U67480; NOT_ANNOTATED_CDS.
DR PIR: A64332; A64332.
DR TIGR: MJ0256;
DR InterPro: IPR000399; Pyruvate decarb.
DR Pfam: PF02775; TPP_enzymes.C.1.
DR PROSITE: PS00187; TPP_ENZYMES; FALSE_NEG.
KW Coenzyme M biosynthesis; Lyase; Decarboxylase; Thiamine pyrophosphate; Complete proteome.
SO SEQUENCE 188 AA; 20980 MM; 52A6C91E2D43B97D CRC64;

Query Match 10.4%; Score 67.5; DB 1; Length 188;
Best Local Similarity 24.8%; Pred. No. 5.4;

Matches 26; Conservative 13; Mismatches 45; Indels 21; Gaps 3;
QY 29 GIGILVILGVLILGICWTCRRNGRYALMDK-----LHVGTQALNRCPQEGF 79
DB 72 GGGSLIMNIGSLTIG--YMNKRYILVLIIDNSAYGSGTGNKTHTKNTLEELAKGCG 129
QY 80 D-----HRDSKVSIOEKNCPEVYPNAPRAYKLEAEQSP 114
DB 130 DTTTSELEEFERKRNLEKCKVIAKTIPIYNEKCSNIEIIP 174

RESULT 13
MGDI_HUMAN
ID MGDI_HUMAN STANDARD; PRT; 778 AA.
AC Q9Y3V3; Q9H352; Q9HRT4; Q9QF36;
DT 16-OCT-2001 (Rel. 40, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Melanoma-associated antigen DI (MAGE-DI antigen) (Neurotrophin receptor-interacting MAGE homology) (PRO22922).
GN MAGE-DI OR NRAGE.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Bone marrow;
RX MEDLINE=99339980; PubMed=10409427;
RA Pold M., Zhou J., Chen G.L., Hall J.M., Vescio R.A., Berenson J.R.;
RT "Identification of a new, unorthodox member of the MAGE gene family."; Genomics 59:161-167(1999).
RL [12]
RN RP SEQUENCE FROM N.A.
RX MEDLINE=20439481; PubMed=10985348;
RA Salehi A.H., Roux P.P., Kubu C.J., Zetindler C., Bhakar A.,
RT "NRAGE, a novel MAGE protein, interacts with the p75 neurotrophin receptor and facilitates nerve growth factor dependent apoptosis."; Neuron 27:279-288(2000).
RL [13]
RN RP SEQUENCE OF 304-778 FROM N.A.
RC TISSUE-Testis;
RA Blum H., Bauersachs S., Mewes H.-W., Gassenhuber J., Wiemann S.;
RT Submitted (DEC-1999) to the EMBL/Genbank/DBJ databases.
RL [4]
RN RP SEQUENCE OF 396-778 FROM N.A.
RC TISSUE=Fetal liver;
RX MEDLINE=21177478; PubMed=11280991;
RA Zhang C.G., Xing G.C., Wei H.D., Yu Y.T., He F.C.;
RT "A new melanoma antigen-encoding gene subfamily in human chromosome X."; J. Clin. Oncol. 19:203-209(2001).
RL [5]
RN RP IDENTIFICATION OF THE TRANSLATIONAL INITIATION CODON.
RX MEDLINE=20541720; PubMed=11087672;
RA Kubu C.J., Goldhawk D.G., Barker P.A., Verdi J.M.;
RT "Identification of the translational initiation codon in human MAGE-DI."; Genomics 70:150-152(2000).
RL [1]
RN RP FUNCTION: Involved in the apoptotic response after nerve growth factor (NGF) binding in neuronal cells. Binds p75NTR and antagonizes its association with TrkA, inhibits cell cycle progression, and facilitates p75NTR-mediated apoptosis. May act as a regulator of the function of DLX family members.
CC -1- SUBUNIT: Interacts with the p75 neurotrophin receptor.
CC -1- SUBCELLULAR LOCATION: Cytoplasmic. Expression shifts from the cytoplasm to the plasma membrane upon stimulation with NGF (By similarity).
CC -1- TISSUE SPECIFICITY: EXPRESSED IN BONE MARROW STROMAL CELLS FROM BOTH MULTIPLE MYELOMA PATIENTS AND HEALTHY DONORS. SEEMS TO BE UBQUITOUSLY EXPRESSED.
CC -1- SIMILARITY: Contains 1 MAGE domain.

CC -1- CAUTION: REF.1 differs from that shown due to several frameshifts that resulted in a N-terminally truncated protein.
CC
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CC
CC EMBL: AF124440; AAC31421.1; ALT_FRAME.
CC EMBL: AF217963; AAC09704.1; -
CC EMBL: AL133628; CAB63752.1; -
CC EMBL: AF132205; AAC35551.1; ALT_INIT.
CC Genew: HGNC:6813; MAGE-DI.
CC MIM: 300224; -
CC CO: GO:0008222; F: tumor antigen; TAS.
CC InterPro: IPR002190; MAGE.
CC Pfam: PF01454; MAGE; 1.
CC PROSITE: PS50838; MAGE; 1.
CC K1 Antigen; Multigene family; Repeat.
CC DOMAIN 296 444
FT FT REPEAT 296 301 1.
FT FT REPEAT 302 307 2.
FT FT REPEAT 308 313 3.
FT FT REPEAT 332 337 4.
FT FT REPEAT 338 343 5.
FT FT REPEAT 344 349 6.
FT FT REPEAT 350 355 7.
FT FT REPEAT 356 361 8.
FT FT REPEAT 362 367 9.
FT FT REPEAT 368 373 10.
FT FT REPEAT 374 379 11.
FT FT REPEAT 380 385 12.
FT FT REPEAT 386 391 13.
FT FT REPEAT 392 397 14.
FT FT REPEAT 398 403 15.
FT FT REPEAT 404 409 16.
FT FT REPEAT 410 415 17.
FT FT REPEAT 416 421 18.
FT FT REPEAT 422 427 19.
FT FT REPEAT 428 432 20 (IMPERFECT).
FT FT REPEAT 433 438 21.
FT FT REPEAT 439 444 22.
FT FT DOMAIN 471 669 MAGE.
SQ SEQUENCE 778 AA; 86150 MW; 0F8BEC7155326FCC CRC64;
Query Match 10.48; Score 67.5; DB 1; Length 778;
Best Local Similarity 29.78; Pred. No. 25;
Matches 33; Conservative 10; Mismatches 33; Indels 35; Gaps 6;
QY 19 HSY---TTAEAGI-----GILVILGVLILG-----CWGCRNGRYALM 58
DB 536 HLYILSTPSLAGILGTTDTPTRLGLVILGVIYFNNGRASEAVLWELARKGLRPGV 595
QY 59 DKSLHVGTCALYTR-CPQGFPHRDSKVSIOEKNCPEVYPNA-PPAYEK 107
DB 596 RHPLDGLRLKLYEFVKQKLYYR-----VPSNPPRYEFL 633
RESULT 14
FATH_HUMAN
ID FATH_HUMAN STANDARD; PRT; 4590 AA.
AC Q14517;
DT 16-OCT-2001 (Rel. 40, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Cadherin-related tumor suppressor precursor (Fat protein homolog).
GN FAT.
OS Homo sapiens (Human).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.
 OX NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Lymphocytes; PubMed=8586420;
 RX MEDLINE=96163873; PubMed=8586420;
 RA Dunne J., Hanby A.M., Poulson R., Jones T.A., Sheer D., Chin W.G.,
 Da S.M., Zhao Q., Beverley P.C.L., Owen M.J.;
 RT "Molecular cloning and tissue expression of FAT, the human homologue
 of the Drosophila fat gene that is located on chromosome 4q34-q35 and
 encodes a putative adhesion molecule.";
 RL Genomics 30:207-223(1995).
 CC -1 FUNCTION: COILED FUNCTION AS A CELL-ADHESION PROTEIN.
 CC -1 SUBCELLULAR LOCATION: Type I membrane protein (By similarity).
 CC -1 TISSUE SPECIFICITY: EXPRESSED IN MANY EPITHELIAL AND SOME
 CC -1 ENDOTHELIAL AND SMOOTH MUSCLE CELLS.
 CC -1 SIMILARITY: Contains 34 cadherin domains.
 CC -1 SIMILARITY: Contains 5 EGF-like domains.
 CC -1 SIMILARITY: Contains 1 laminin G-like domain.
 CC -----
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 CC -----
 DR EMBL; X87241; CA60685.1; -
 DR HSSP; P01132; IEGF.
 DR Genew; HGNC:3595; FAT.
 DR MIM; 600976; -
 DR GO; GO:0005887; C: Integral to plasma membrane; TAS.
 DR GO; GO:0008181; F: tumor suppressor; TAS.
 DR GO; GO:0007155; P: cell adhesion; TAS.
 DR GO; GO:0007267; P: cell-cell signaling; TAS.
 DR GO; GO:0007345; P: embryogenesis and morphogenesis; TAS.
 DR InterPro; IPR000152; Asx_hydroxyl.
 DR InterPro; IPR002126; Cadherin.
 DR InterPro; IPR000742; EGF_2.
 DR InterPro; IPR001881; EGF_Ca.
 DR InterPro; IPR006209; EGF_like.
 DR InterPro; IPR001791; Laminin_G.
 DR Pfam; PF000028; cadherin; 33.
 DR Pfam; PF00008; EGF; 5.
 DR Pfam; PF00054; Laminin_G; 1.
 DR PRINTS; PR00205; CADHERIN.
 DR SMART; SM00112; CA; 31.
 DR SMART; SM00179; EGF_CA; 1.
 DR SMART; SM00282; Lamg; 1.
 DR PROSITE; PS00010; ASX_HYDROXYL; 1.
 DR PROSITE; PS00232; CADHERIN_1; 16.
 DR PROSITE; PS0268; CADHERIN_2; 33.
 DR PROSITE; PS00022; EGF_1; 4.
 DR PROSITE; PS01186; EGF_2; 1.
 DR PROSITE; PS01187; EGF_CA; 1.
 DR PROSITE; PS50025; LAM_G_DOMAIN; 1.
 DR Cell adhesion; Signal; Glycoprotein; Transmembrane; Calcium-binding;
 KW Repeat; EGF-like domain.
 FT SIGNAL 1 21
 FT CHAIN 22 4590
 FT FT
 FT DOMAIN 22 4183
 FT TRANSMEM 4184 4204
 FT DOMAIN 4205 4590
 FT DOMAIN 22 149
 FT DOMAIN 150 256
 FT DOMAIN 257 361
 FT DOMAIN 362 463
 FT DOMAIN 464 569
 FT DOMAIN 570 716
 FT DOMAIN 717 822
 FT DOMAIN 717 822

FT DOMAIN 823 927
 FT DOMAIN 928 1034
 FT DOMAIN 1035 1138
 FT DOMAIN 1139 1245
 FT DOMAIN 1246 1345
 FT DOMAIN 1346 1456
 FT DOMAIN 1457 1562
 FT DOMAIN 1563 1670
 FT DOMAIN 1671 1769
 FT DOMAIN 1770 1882
 FT DOMAIN 1883 1982
 FT DOMAIN 1983 2084
 FT DOMAIN 2085 2185
 FT DOMAIN 2186 2286
 FT DOMAIN 2287 2393
 FT DOMAIN 2394 2495
 FT DOMAIN 2496 2599
 FT DOMAIN 2600 2705
 FT DOMAIN 2706 2811
 FT DOMAIN 2812 2920
 FT DOMAIN 2921 3024
 FT DOMAIN 3025 3127
 FT DOMAIN 3128 3232
 FT DOMAIN 3233 3337
 FT DOMAIN 3338 3442
 FT DOMAIN 3443 3546
 FT DOMAIN 3547 3649
 FT DOMAIN 3649 3827
 FT DOMAIN 3831 4011
 FT DOMAIN 4013 4051
 FT DOMAIN 4052 4085
 FT DOMAIN 4090 4126
 FT DOMAIN 4127 4163
 FT DOMAIN 4163 40
 FT CARBOHYD 40 40
 FT CARBOHYD 333 333
 FT CARBOHYD 660 660
 FT CARBOHYD 740 740
 FT CARBOHYD 791 791
 FT CARBOHYD 998 998
 FT CARBOHYD 1426 1426
 FT CARBOHYD 1551 1551
 FT CARBOHYD 1751 1751
 FT CARBOHYD 1867 1867
 FT CARBOHYD 1905 1905
 FT CARBOHYD 1943 1943
 FT CARBOHYD 1994 1994
 FT CARBOHYD 2328 2328
 FT CARBOHYD 2467 2467
 FT CARBOHYD 3326 3326
 FT CARBOHYD 3424 3424
 FT CARBOHYD 3446 3446
 FT CARBOHYD 3615 3615
 FT CARBOHYD 3642 3642
 FT CARBOHYD 3718 3718
 FT CARBOHYD 4154 4154
 FT SEQUENCE 4590 AA; 506273 MW; 04483CCDD00860A7 CRC64;
 Query Match 10.4%; Score 67.5; DB 1; Length 4590;
 Best Local Similarity 27.7%; Pred. No. 1.6e+02;
 Matches 31; Conservative 14; Mismatches 38; Indels 29; Gaps 6;
 OY 27 AAGTGLVITLVLLIGCW-YCR---RNGYRALDKSLVGTQCALRRCPQSGFDR 82
 DB 4182 AAGTGLVITLVLLIGCW-YCR---RNGYRALDKSLVGTQCALRRCPQSGFDR 82
 OY 83 DSKVSGJOEKN---CEPVVFNAPVY-----EKLSSAOSPPRSP 118
 DB 4236 DSKLN---KNITSDIPVPPVPISTTSPISDSRNNDNSPFGSAIPBHP 4284
 RESULT 15
 MGD1_MOUSE
 ID MGD1_MOUSE STANDARD; PRT; 775 AA.

AC 090YH6; 090PB5; 09CYX1;
 DT 28-FEB-2003 (Rel. 41, Created)
 DT 28-FEB-2003 (Rel. 41, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Melanoma-associated antigen DI (MAGE-DI antigen) (Neurotrophin
 receptor-interacting MAGE homolog) (Dlx1n-1).
 GN MAGE-DI OR NRAGE.
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 NC NCBL:taxid:10090;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Embryo;
 RX MEDLINE=21265065; PubMed=11084035;
 RA Masuda Y., Sasaki A., Shibuya H., Ueno N., Ikeda K., Watanabe K.;
 RT Dlx1n-1, a novel protein that binds Dlx5 and regulates its
 RT transcriptional function.";
 RL J. Biol. Chem. 276:5331-5338(2001).
 RN [2]
 RP SEQUENCE FROM N.A.
 RA Auguter P.H., Chomez P.M., De Backer O.R., Bertrand M.J.M.;
 RT "Ten new murine members of the MAGE gene family.";
 RL Submitted (NOV-2000) to the EMBL/Genbank/DBJ databases.
 RN [3]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=Head;
 RX MEDLINE=21085660; PubMed=11217851;
 RA Kawai J., Shinagawa A., Shibata K., Yoshino M., Itoh M., Ishii Y.,
 RA Aizawa T., Hara A., Fukunishi Y., Konno H., Adachi J., Fukuda S.,
 RA Aizawa K., Izawa M., Nishi K., Kiyosawa H., Kondo S., Yamana K.I.,
 RA Saito T., Okazaki Y., Gojobori T., Bono H., Kasukawa T., Saito R.,
 RA Kadota K., Matsuda H.A., Ashburner M., Batalov S., Casavant T.,
 RA Fleischmann W., Gaasterland T., Glass C., King B., Kochiya H.,
 RA Kuehl P., Lewis S., Matsuo Y., Nikaido I., Pesole G., Quackenbush J.,
 RA Schiml L.M., Stabili F., Suzuki R., Tomita M., Wagner L., Washio T.,
 RA Sakai K., Okido T., Furuno M., Aono H., Baldarelli R., Barsh G.,
 RA Blake J., Boftelli D., Bojunga N., Carninci P., de Bonaldo M.F.,
 RA Brownstein M.J., Bult C., Fletcher C., Fujita M., Gariboldi M.,
 RA Guslinich S., Hill D., Hofmann M., Hume D.A., Kamaya M., Lee N.H.,
 RA Lyons P., Marchionni L., Mashima J., Mazzarelli J., Mombaerts P.,
 RA Morone P., Ring B., Ringwald M., Rodriguez I., Sakamoto N.,
 RA Sasaki H., Sato K., Schoenbach C., Seya T., Shibata Y., Storch K.-F.,
 RA Suzuki H., Toyooka K., Wang K.H., Weitz C., Whitaker C., Wilming L.,
 RA Wyshak-Boris A., Yoshida K., Hasegawa Y., Kawaji H., Kohsaki S.,
 RA Hayashizaki Y.;
 RT *Functional annotation of a full-length mouse cDNA collection.";
 RL Nature 403:685-690(2001).
 CC -1- FUNCTION: Involved in the apoptotic response after nerve growth
 CC factor (NGF) binding in neuronal cells. Binds p75NTR and
 CC antagonizes its association with TrkA, inhibits cell cycle
 CC progression, and facilitates p75NTR-mediated apoptosis. May act as
 CC a regulator of the function of Dlx family members (By similarity).
 CC -1- SUBUNIT: INTERACTS WITH Dlx5, Dlx7 AND MSX2 AND FORMS
 CC HOMOMULTIMERS.
 CC -1- SUBCELLULAR LOCATION: Cytoplasmic. Expression shifts from the
 CC cytoplasm to the plasma membrane upon stimulation with NGF (By
 CC similarity).
 CC -1- TISSUE SPECIFICITY: Ubiquitously expressed in many adult tissues,
 CC except for the spleen. Expressed in osteoblastic and
 CC chondrogenic cell lines and also during embryonic development.
 CC -1- SIMILARITY: Contains 1 MAGE domain.
 CC -----
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
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 CC -----
 CC DR EMBL; AB029448; BAA87959.1; -;
 CC DR EMBL; AF319975; AAK01203.1; -;

DR EMBL; AK017275; BAB30666.1; -;
 DR EMBL; AK013231; -; NOT_ANNOTATED_CDS.
 DR MGD; MGI:1930187; Maged1.
 DR GO; GO:0005515; F:protein binding activity; IPI.
 DR GO; GO:0003713; F:transcription co-activator activity; IDA.
 DR GO; GO:0006357; P:regulation of transcription from Pol II pro. . .; IDA.
 DR InterPro; IPR002190; MAGE.
 DR Pfam; PF01454; MAGE; 1.
 DR PROSITE; PS50838; MAGE; 1.
 KW Antigen; Multigene family; Repeat.
 FT DOMAIN 292 441
 FT REPEAT 292 297 X.
 FT REPEAT 298 303 1.
 FT REPEAT 304 309 2.
 FT REPEAT 329 334 3.
 FT REPEAT 335 340 4.
 FT REPEAT 341 346 5.
 FT REPEAT 347 352 6.
 FT REPEAT 353 358 7.
 FT REPEAT 359 364 8.
 FT REPEAT 365 370 9.
 FT REPEAT 371 376 10.
 FT REPEAT 377 382 11.
 FT REPEAT 383 388 12.
 FT REPEAT 389 394 13.
 FT REPEAT 395 400 14.
 FT REPEAT 401 406 15.
 FT REPEAT 407 412 16.
 FT REPEAT 413 418 17.
 FT REPEAT 419 424 18.
 FT REPEAT 425 429 19.
 FT REPEAT 430 435 20 (IMPERFECT).
 FT REPEAT 436 441 21.
 FT REPEAT 448 466 22.
 FT DOMAIN 357 666 MAGE.
 FT CONFLICT 357 662 MISSING (IN REF. 2).
 SQ SEQUENCE 775 AA; 85669 MW; 224B82470816835A CRC64;
 Query Match 10.2%; Score 66.5; DB 1; Length 775;
 Best Local Similarity 28.8%; Pred. No. 31;
 Matches 32; Conservative 11; Mismatches 33; Indels 35; Gaps 6;
 QY 19 HSY---TTAEBAAGI-----GIIVTIVGLILIG-----CWYCRNRNGYRALM 58
 DB 533 HLYVLISTPESLNGILGTTKDTPTLGLILVILGIIFPNNGNATNAVLEMLRRKGLRGV 592.
 QY 59 DKSLVGTQCALRR-CPQEGFDHRSKVSLSQENCEPVVPA-PRAVEKL 107
 DB 593 RHPILGDLRLKLLTYEFVKOKLYDRR-----VNSNPPEVEFL 630
 Search completed: October 7, 2003, 18:48:34
 Job time : 13 secs